

GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: July 27, 2004, 17:59:42 ; Search time 2777 Seconds  
(without alignments)  
2419.514 Million cell updates/sec

Title: US-09-765-231A-58  
Perfect score: 225  
Sequence: 1 tgaaggtaagtgttcagg.....attaggaattttttttttt 225

Scoring table: IDENTITY NUC  
Gapop 10.0 , Gapext 1.0

Searched: 27513289 seqs, 14931090276 residues  
Total number of hits satisfying chosen parameters: 55026578

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000  
Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database :

EST:\*

1: em\_estba:\*  
2: em\_esthum:\*  
3: em\_estin:\*  
4: em\_estmu:\*  
5: em\_estov:\*  
6: em\_estpl:\*  
7: em\_estro:\*  
8: em\_estl:\*  
9: gb\_est1:\*  
10: gb\_est2:\*  
11: gb\_est3:\*  
12: gb\_est4:\*  
13: gb\_est5:\*  
14: gb\_est6:\*  
15: em\_estfun:\*  
16: em\_estom:\*  
17: em\_gss\_hum:\*  
18: em\_gss\_inv:\*  
19: em\_gss\_pln:\*  
20: em\_gss\_vrt:\*  
21: em\_gss\_fun:\*  
22: em\_gss\_mam:\*  
23: em\_gss\_mus:\*  
24: em\_gss\_pro:\*  
25: em\_gss\_rod:\*  
26: em\_gss\_phg:\*  
27: em\_gss\_vrl:\*  
28: gb\_gss1:\*  
29: gb\_gss2:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	196	87.1	341	9	AA502552 ng62e06.s
2	117	52.0	324	9	AI247782 qh61a07.x
3	117	52.0	362	9	AA682512 zii9a01.s
4	48.8	21.7	928	29	CNS00DKY AL071865 Drosophila

C	5	47	20.9	1131	13	BX356147
C	6	44.6	19.8	1201	13	BX324729
C	7	44.2	19.6	1108	13	BX418757
C	8	43.8	19.5	1155	9	AL514851
C	9	43	19.1	773	28	AQ781761
C	10	42.8	19.0	1091	13	BX424950
C	11	42.4	18.8	1121	13	BX338325
C	12	42.2	18.8	542	13	BUT22265
C	13	42.2	18.8	1056	13	BM415058
C	14	42	18.7	1201	13	BM462207
C	15	41.8	18.6	964	29	CNS058MA
C	16	41.2	18.3	1101	29	CNS00D90
C	17	41	18.2	290	10	AM504318
C	18	41	18.2	451	14	CB051834
C	19	41	18.2	1562	11	BC022863
C	20	40.6	18.0	595	28	AZ523166
C	21	40.6	18.0	957	29	CNS015W7
C	22	40.4	18.0	516	13	BM561821
C	23	40.4	18.0	525	10	AW381028
C	24	40.4	18.0	999	13	BM380865
C	25	40.4	18.0	1201	13	BM385531
C	26	40	17.8	475	10	BE325739
C	27	40	17.8	611	29	CE160147
C	28	40	17.8	945	13	BM418213
C	29	40	17.8	1081	28	CC247576
C	30	40	17.8	1098	13	BM377526
C	31	40	17.8	1201	13	BM458169
C	32	39.8	17.7	208	9	AU071524
C	33	39.8	17.7	638	14	CD649691
C	34	39.8	17.7	647	28	B83740
C	35	39.8	17.7	800	14	CB971606
C	36	39.8	17.7	890	14	CB756780
C	37	39.8	17.7	1201	13	BM421216
C	38	39.6	17.6	358	14	CD801440
C	39	39.6	17.6	360	9	AL750985
C	40	39.6	17.6	409	13	BQ451684
C	41	39.6	17.6	524	28	AZ055107
C	42	39.6	17.6	941	28	BH133309
C	43	39.6	17.6	1195	13	BM355698
C	44	39.4	17.5	189	12	BJ351968
C	45	39.4	17.5	985	9	AL520226

ALIGNMENTS

RESULT 1  
LOCUS AA502552 341 bp mRNA linear EST 19-AUG-1997  
DEFINITION ng62e06.s1 NCI\_CGAP\_Lip2 Homo sapiens cDNA clone IMAGE:939394, mRNA sequence.  
ACCESSION AA502552  
VERSION AA502552.1 GI:2237519  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
REFERENCE 1 (bases 1 to 341)  
AUTHORS NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.  
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index  
JOURNAL Unpublished (1997)  
COMMENT Contact: Robert Strausberg, Ph.D.  
Email: [cgaps-r@mail.nih.gov](mailto:cgaps-r@mail.nih.gov)  
Tissue Procurement: L. Jeffrey Medeiros, M.D., Michael R. Emmer-Buck, M.D., Ph.D.  
cDNA Library Preparation: David B. Krizman, Ph.D.  
DNA Sequencing by: Washington University Genome Sequencing Center  
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: [www-bio.llnl.gov/bbrp/image/image.html](http://www-bio.llnl.gov/bbrp/image/image.html)

Insert Length: 1183 Std Error: 0.00  
Seq primer: -40ml3 fwd. ET from Amersham  
High quality sequence stop: 310.

#### FEATURES

Location/Qualifiers  
1..341  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="IMAGE:939394"  
/tissue\_type="liposarcoma"  
/lab\_host="DH10B"  
/clone\_lib="NCI-CGAP\_Lip2"  
/note="Vector: pAMP10; mRNA made from liposarcoma, cDNA made by oligo-dT priming. Non-directionally cloned. Size-selected on agarose gel, average insert size 600 bp. Reference: Krizman et al. (1996) Cancer Research 56:5380-5383."

#### ORIGIN

Query Match 87.1%; Score 196; DB 9; Length 341;  
Best Local Similarity 99.5%; Pred. No. 5.9e-28;  
Matches 207; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY 19 GGCATAAAATTGAAATTAATATGAGGCTCATGATGCTATATGTTTACCTTCA 78  
Db |||||||  
8 GGCATAAAATTGAAATTAATATGAGGCTCATGATGCTATATGTTTACCTTCA 67  
QY 79 GAAGATAATTAGTTTCACTCAGGTTTTTCAAGCTACGCTGCCCAAAAACGAAC 138  
Db |||||||  
68 GAAGATAATTAGTTTCACTCAGGTTTTTCAAGCTACGCTGCCCAAAAACGAAC 127  
QY 139 AAAAC-AAAAAACAACCTTTTAAAGTTGATGCTACTCATTTGATCGCTCCTCTG 197  
Db |||||||  
128 AAAACAAAAAACAACCTTTTAAAGTTGATGCTACTCATTTGATCGCTCCTCTG 187  
QY 198 CTGAATCAATTAGGAATTTTTTTTTT 225  
Db |||||||  
188 CTGAATCAATTAGGAATTTTTTTTTT 215

RESULT 2  
LOCUS AI247782 324 bp mRNA linear EST 01-DEC-1998  
DEFINITION Q61a07.x1 Soares\_fetal\_liver\_spleen\_INFLS\_S1 Homo sapiens cDNA  
clone IMAGE:1849140 3', mRNA sequence.  
ACCESSION AI247782  
VERSION AI247782.1 GI:3843179  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
REFERENCE 1 (bases 1 to 324)  
AUTHORS NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.  
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index  
JOURNAL Unpublished (1997)  
COMMENT Contact: Robert Strausberg, Ph.D.  
Email: [cgaps-remail.nih.gov](mailto:cgaps-remail.nih.gov)  
This clone is available royalty-free through LLNL ; contact the IMAGE Consortium ([infoimage.llnl.gov](http://infoimage.llnl.gov)) for further information.  
Insert Length: 821 Std Error: 0.00  
Seq primer: -40UP from Gibco  
High quality sequence stop: 317.

Location/Qualifiers  
1..324  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="IMAGE:1849140"  
/sex="male"  
/dev\_stage="20 week-post conception fetus"  
/lab\_host="DH10B (ampicillin resistant)"

#### FEATURES

Location/Qualifiers  
1..324  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="IMAGE:1849140"  
/sex="male"  
/dev\_stage="20 week-post conception fetus"  
/lab\_host="DH10B (ampicillin resistant)"

/clone\_lib="Soares fetal liver spleen\_INFLS\_S1"  
/note="Organ: Liver and Spleen; Vector: p7773D (Pharmacia) with a modified polylinker; Site 1: Pac I; Site 2: Eco RI; This is a subcloned version of the original Soares fetal liver spleen INFLS library. 1st strand cDNA was primed with a Pac I - oligo(dT) primer [5', AACTGGAAGAATTAAATAAGATCTTTTTTTTTTTT 3'], double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Pac I and cloned into the Pac I and Eco RI sites of the modified p7773 vector. Library went through one round of normalization. Library constructed by Bento Soares and M.Fatima Bonaldo."

#### ORIGIN

Query Match 52.0%; Score 117; DB 9; Length 324;  
Best Local Similarity 100.0%; Pred. No. 7e-13;  
Matches 117; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 109 AAAGCTACGCTGCCCAAAAACGAAACAAAACAAAACAAACACCTTTTAAAGATTG 168  
Db |||||||  
1 AAAGCTACGCTGCCCAAAAACGAAACAAAACAAAACAAACACCTTTTAAAGATTG 60  
QY 169 ATGCTACTCATTTGATCTGCTCTCTGCTCAATCAATTAGGAATTTTTTTTTT 225  
Db |||||||  
61 ATGCTACTCATTTGATCTGCTCTCTGCTCAATCAATTAGGAATTTTTTTTTT 117

RESULT 3  
LOCUS AA682512 362 bp mRNA linear EST 19-DEC-1997  
DEFINITION z119a01.s1 Soares\_fetal\_liver\_spleen\_INFLS\_S1 Homo sapiens cDNA  
clone IMAGE:431208 3', mRNA sequence.

ACCESSION AA682512  
VERSION AA682512.1 GI:2669793  
KEYWORDS EST.

SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens

Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
REFERENCE 1 (bases 1 to 362)  
AUTHORS Hallier, L., Allen, M., Bowles, L., Dubuque, T., Geisel, G., Jost, S., Krizman, D., Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin, J., Moore, B., Schellenberg, K., Steptoe, M., Tan, F., Theising, B., White, Y., Wyllie, T., Waterston, R. and Wilson, R.

WashU-NCI human EST Project  
Unpublished (1997)  
Contact: Wilson RK

Washington University School of Medicine  
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
Tel: 314 286 1800  
Fax: 314 286 1810  
Email: [est@wustl.edu](mailto:est@wustl.edu)

This clone is available royalty-free through LLNL ; contact the IMAGE Consortium ([infoimage.llnl.gov](http://infoimage.llnl.gov)) for further information.

Seq primer: -40ml3 fwd. ET from Amersham

High quality sequence stop: 308.

Location/Qualifiers  
1..362  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="IMAGE:431208"  
/sex="male"  
/dev\_stage="20 week-post conception fetus"  
/lab\_host="DH10B (ampicillin resistant)"

/clone\_lib="Soares fetal liver spleen\_INFLS\_S1"  
/note="Organ: Liver and Spleen; Vector: p7773D (Pharmacia) with a modified polylinker; Site 1: Pac I; Site 2: Eco RI; This is a subcloned version of the original Soares fetal liver spleen INFLS library. 1st strand cDNA was primed with a Pac I - oligo(dT) primer [5', AACTGGAAGAATTAAATAAGATCTTTTTTTTTTTT 3'], double-stranded cDNA was ligated to Eco RI adaptors

(Pharmacacia), digested with Pac I and cloned into the Pac I and Eco RI sites of the modified pT73 vector. Library went through one round of normalization. Library constructed by Bento Soares and M.Fatima Bonaldo."

## ORIGIN

Query Match 52.0%; Score 117; DB 9; Length 362;  
Best Local Similarity 100.0%; Pred. No. 6.5e-13;  
Matches 117; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 109 AAGCTACGCTGCCCAAAACGAACAAACAAAACAAACAAACCTTTTAAAGAGTTG 168

Db 1 AAGCTACGCTGCCCAAAACGAACAAACAAAACAAACAAACCTTTTAAAGAGTTG 60

Qy 169 ATGGCTACTCATTTGATCTGCTCTGCTGATCAATTAGGAATTTTTTTTTT 225

Db 61 ATGGCTACTCATTTGATCTGCTCTGCTGATCAATTAGGAATTTTTTTTTT 117

## RESULT 4

CNS00DKY 928 bp DNA linear GSS 04-JUN-1999  
LOCUS Drosophila melanogaster genome survey sequence T7 end of BAC #  
DEFINITION BACR27A24 of RPCI-98 library from Drosophila melanogaster (fruit fly), genomic survey sequence.

ACCESSION AL071865

VERSION AL071865.1 GI:4948170

SOURCE GSS.

ORGANISM Drosophila melanogaster (fruit fly)

Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.  
1 (bases 1 to 928)

## REFERENCE

AUTHORS

TITLE

JOURNAL

## COMMENT

Direct Submission  
Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage :  
BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr  
- Web : www.genoscope.cns.fr)  
Determination of this BAC-end sequence was carried out as part of a collaboration with the Berkeley Drosophila Genome Project (BDGP). The BDGP is constructing a physical map of the Drosophila melanogaster genome using these BACs. For further information please see <http://www.fruitfly.org> The BDGP Drosophila melanogaster BAC library was prepared by Kazutoyo Osogawa and Aaron Mamoser in Pieter de Jong's laboratory in the Department of Cancer Genetics at the Roswell Park Cancer Institute in Buffalo, NY. The library is named RPCI-98 and was constructed by partial EcoRI digestion of Drosophila DNA provided by the BDGP from the isogenic strain Y2; cn bw sp, the same strain used for the BDGP's P1 and EST libraries. A more detailed description of the library and how to order individual BAC clones, the entire library, or filters for hybridization from the BACPAC Resource Center can be found at [http://bacpac.med.buffalo.edu/drosophila\\_bac.htm](http://bacpac.med.buffalo.edu/drosophila_bac.htm).

## FEATURES

source

1..928

/organism="Drosophila melanogaster"

/mol\_type="genomic DNA"

/db\_xref="taxon:7227"

/clone="BACR27A24"

/clone\_lib="RPCI-98"

/note="end : T7"

Query Match 21.7%; Score 48.8; DB 29; Length 928;

Best Local Similarity 31.4%; Pred. No. 3.5;  
Matches 64; Conservative 59; Mismatches 81; Indels 0; Gaps 0;

Qy 22 ATAAATTTGAATAAATATGAGGCTCCATGATGCTATATGTTTACCTTCAGAA 81

Db 678 AAAAATTTAATAAATAAATAAATTAATTTTTTTTTTTTATATWATAWAAA 737

Qy 82 GAATATTAGTTTCACTCAGGTTTTTCAAGCTACGCTGCCCAAAACGAAACAA 141

Db 738 TATWWWWATWTWDGKNWNNNAWWWWWWWWWWWWWWWWWWWWWWWWWWWW 797  
Qy 142 ACAAAAAACACCTTTTAAAGAGTTGATGGCTACTCATTTGATCTGCTCTGTA 201  
Db 798 AAAAATAAATAAATATGAGGCTCCATGATGCTCCCTCTCTGTA 857  
Qy 202 ATCAATTAGGAATTTTTTTTTT 225  
Db 858 RWWTTTTTTTTTTTTTTTTTTT 881

## RESULT 5

EX356147/c

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

1..1131

/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/clone="CSODI008YB06"

/tissue\_type="PLACENTA"

/note="1st strand cDNA was primed with a NotI-oligo (dT) primer. Five prime end enriched, double-strand cDNA was digested with NotI and cloned into the NotI and EcoR V sites of the pCMVSPORT 6 vector. Library was normalized."

Location/Qualifiers

1..1131

/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/clone="CSODI008YB06"

/tissue\_type="PLACENTA"

/note="1st strand cDNA was primed with a NotI-oligo (dT) primer. Five prime end enriched, double-strand cDNA was digested with NotI and cloned into the NotI and EcoR V sites of the pCMVSPORT 6 vector. Library was normalized."

Location/Qualifiers

1..1131

/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/clone="CSODI008YB06"

/tissue\_type="PLACENTA"

/note="1st strand cDNA was primed with a NotI-oligo (dT) primer. Five prime end enriched, double-strand cDNA was digested with NotI and cloned into the NotI and EcoR V sites of the pCMVSPORT 6 vector. Library was normalized."

Location/Qualifiers

1..1131

/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/clone="CSODI008YB06"

/tissue\_type="PLACENTA"

/note="1st strand cDNA was primed with a NotI-oligo (dT) primer. Five prime end enriched, double-strand cDNA was digested with NotI and cloned into the NotI and EcoR V sites of the pCMVSPORT 6 vector. Library was normalized."

Location/Qualifiers

1..1131

/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/clone="CSODI008YB06"

/tissue\_type="PLACENTA"

/note="1st strand cDNA was primed with a NotI-oligo (dT) primer. Five prime end enriched, double-strand cDNA was digested with NotI and cloned into the NotI and EcoR V sites of the pCMVSPORT 6 vector. Library was normalized."

Location/Qualifiers

1..1131

/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/clone="CSODI008YB06"

/tissue\_type="PLACENTA"

/note="1st strand cDNA was primed with a NotI-oligo (dT) primer. Five prime end enriched, double-strand cDNA was digested with NotI and cloned into the NotI and EcoR V sites of the pCMVSPORT 6 vector. Library was normalized."

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RESULT 6
LOCUS      BX324729/c
DEFINITION BX324729 Homo sapiens PLACENTA COT 25-NORMALIZED Homo sapiens cDNA
            clone CS0D1037YJ05 5-PRIME, mRNA sequence.
ACCESSION  BX324729
VERSION     BX324729.1 GI:30338394
KEYWORDS   EST.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
REFERENCE  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
AUTHORS    Li,W.B., Gruber,C., Jessee,J. and Polayes,D.
TITLE      Full-length cDNA libraries and normalization
JOURNAL    Unpublished (2001)
COMMENT    Contact: Genoscope
            Genoscope - Centre National de Sequencage
            BP 191 91006 EVRY cedex - France
            Email: seqref@genoscope.cns.fr, Web : www.genoscope.cns.fr
            Library was constructed by Life Technologies, a division of
            Invitrogen. Contact : Feng Liang Email : fliang@lifetech.com URL :
            http://fulllength.invitrogen.com/ Invitrogen Corporation 1600
            Faraday Avenue Genoscope sequence ID : CS0D1037CE03QPI.
            Location/Qualifiers
                1..1201
                /organism="Homo sapiens"
                /mol_type="mRNA"
                /db_xref="taxon:9606"
                /clone="CS0D1037YJ05"
                /tissue_type="PLACENTA COT 25-NORMALIZED"
                /note="1st strand cDNA was primed with a NotI-oligo(dT)
                primer. Five prime end enriched, double-strand cDNA was
                digested with Not I and EcoR V
                sites of the pCMVSPORT 6 vector. Library was normalized."
FEATURES   source
            1..1201
            Query Match      19.8%; Score 44.6; DB 13; Length 1201;
            Best Local Similarity 36.3%; Pred. No. 19;
            Matches 81; Conservative 41; Mismatches 101; Indels 0; Gaps 0;

QY 3 ATGGTAAGTTGTTTCAGGCATATAAATTGAAATAAATTATGAGGCTCCATGATATGCTAT 62
Db  ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
695 AAGRGAWRGKKTCTMAGGATMTAVGGTGAATAAAAMSYCAARSCTRGCTTAAAAAA 636
QY 63 ATTGGTTTTACCTTCAGAGATATTTAGTTTCTACTCAGGTTTTTCAAGCTACGCTGC 122
Db  ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
635 AAAAAATAATAAAAAAATATATAAATAAAAAAATAAAAAAATAAAAAAATAGGCCCT 576
QY 123 CCCCAAAAAACGAAACAAAAACAAACCTTTTAAAGATTGATGGCTACTCATTT 182
Db  ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
575 CAAATWAAATAAAAAAATAAAAAAATAAAAAAATAAAAAAATAAAAAAATAAAAAA 516
QY 183 GATCGCTCTGCTGCTCAATCAATAGGAATTTTTTTTTTTTTTTT 225
Db  ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
515 TTTWTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT 473

RESULT 7
LOCUS      BX418757/c
DEFINITION BX418757 Homo sapiens FETAL BRAIN Homo sapiens cDNA clone
            CS0DF009YG18 5-PRIME, mRNA sequence.
ACCESSION  BX418757
VERSION     BX418757.1 GI:30769508
KEYWORDS   EST.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
REFERENCE  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
AUTHORS    Li,W.B., Gruber,C., Jessee,J. and Polayes,D.
TITLE      Full-length cDNA libraries and normalization
JOURNAL    Unpublished (2001)
COMMENT    Contact: Genoscope
            Genoscope - Centre National de Sequencage
            BP 191 91006 EVRY cedex - France
            Email: seqref@genoscope.cns.fr, Web : www.genoscope.cns.fr
            Library was constructed by Life Technologies, a division of
            Invitrogen. Contact : Feng Liang Email : fliang@lifetech.com URL :
            http://fulllength.invitrogen.com/ Invitrogen Corporation 1600
            Faraday Avenue Genoscope sequence ID : CS0AI037CE03QPI.
            Location/Qualifiers
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                /mol_type="mRNA"
                /db_xref="taxon:9606"
                /clone="CS0DF009YG18"
                /tissue_type="FETAL BRAIN"
                /dev_stage="fetal"
                /clone_lib="Homo sapiens FETAL BRAIN"
                /note="Torgan; brain; Vector: pCMVSPORT 6; 1st strand cDNA
                was primed with a NotI-oligo(dT) primer. Five prime end
                enriched, double-strand cDNA was digested with Not I and
                cloned into the Not I and EcoRV sites of the pCMVSPORT 6
                vector. Library was not normalized."
FEATURES   source
            1..1108
            Query Match      19.6%; Score 44.2; DB 13; Length 1108;
            Best Local Similarity 32.7%; Pred. No. 23;
            Matches 64; Conservative 49; Mismatches 83; Indels 0; Gaps 0;

QY 30 TGAATAAATATATGAGGCTCCATGATATGCTATATGTTTACCTTCAGAGATATTT 89
Db  ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
722 TCMAYACATTTTCATAATAAATATATATATATATATATATATATATATATATATAT 663
QY 90 AGTTTCTACAGGTTTTTCAAGCTACGCTGTCGCCCAAAAAACGAAACAAAAA 149
Db  ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
662 ATATATNCCAAATTTTTTATATATATAATATATATATATATATATATATATATATAT 603
QY 150 ACRACCTTTTAAAGATTGATGGCTACTCATTTGATGCTGCTCTGCTGAATCAATA 209
Db  ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
602 MCCCMMAAMMMCMCMCMCMCMCMCMCMCMCMCMCMCMCMCMCMCMCMCMCMCMCM 543
QY 210 GGAATTTTTTTTTTTTTT 225
Db  ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
542 WWWWWATTTTTTTTTT 527

RESULT 8
LOCUS      AL514851
DEFINITION AL514851 Homo sapiens NEUROBLASTOMA Homo sapiens cDNA clone
            CL0B8014ZC07 3-PRIME, mRNA sequence.
ACCESSION  AL514851
VERSION     AL514851.2 GI:30464736
KEYWORDS   EST.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
REFERENCE  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
AUTHORS    Li,W.B., Gruber,C., Jessee,J. and Polayes,D.
TITLE      Full-length cDNA libraries and normalization
JOURNAL    Unpublished (2001)
COMMENT    On Feb 13, 2001 this sequence version replaced gi:12778344.
            Contact: Genoscope
            Genoscope - Centre National de Sequencage
            BP 191 91006 EVRY cedex - France

```



Qy

```

REFERENCE
AUTHORS Li,W.B., Gruber,C., Jessee,J. and Polayes,D.
TITLE Full-length cDNA libraries and normalization
JOURNAL Unpublished (2001)
COMMENT Contact: Genoscope
Genoscope - Centre National de Sequencage
BP 191 91006 EVRY cedex - France
Email: seqref@genoscope.cns.fr, Web : www.genoscope.cns.fr
Library was constructed by Life Technologies, a division of
Invitrogen. Contact : Feng Liang Email : fliang@lifetech.com URL :
http://fulllength.invitrogen.com/ Invitrogen Corporation 1600
Faraday Avenue Genoscope sequence ID : CSOCAP004AD10NP1.

FEATURES
source
1. .1056
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="CSOCAP004YG19"
/tissue_type="THYMUS"
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/note="vector: pCMVSPORT 6; 1st strand cDNA was primed
with a NotI-oligo(dT) primer. Five prime end enriched,
double-strand cDNA was digested with Not I and cloned into
the Not I and EcoRV sites of the pCMVSPORT 6 vector.
Library was not normalized."

ORIGIN
Query Match 18.8%; Score 42.2; DB 13; Length 1056;
Best Local Similarity 32.6%; Pred. No. 58;
Matches 70; Conservative 48; Mismatches 97; Indels 0; Gaps 0;

QY 11 TTGTTTCAGGCATGAATTTGAATAAATTATGAGGCTCCATGATGCTATATGCTTT 70
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Db 941 WTTTITTTTNNWWTTTWWAAWTTTWTAAWTTTITTTTITTTTITTTTITTTT 882

QY 71 TACCTTCAGAGATATTAGTTTCACTCAGGTTTTTCAAGCTACGCTGCCCCAAA 130
: ||| : : : : : : : : : : : : : : : : : : : : : : :
Db 881 TWWTTTATAAAWTTAAWAAATTTTITTTTWWTTWWTTWWTTWWTTWWTTWW 822

QY 131 AAGCAACAAACAAACAAACAAACACCTTTTAAAGAGTTGATGGCTACTCATTTGATCTGCC 190
: ||| ||| ||| ||| ||| : : : : : : : : : : : : : : : : :
Db 821 AAAAAAHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHH 762

QY 191 TCCTCTGCTGAATCAATAGAAATTTTITTTTITTTTITTTTITTTTITTTT 225
: ||| : : : : : : : : : : : : : : : : : : : : : : :
Db 761 NATTTTTTTTTTWWAAAAAAWTTTITTTTITTTTITTTTITTTTITTTT 727

RESULT 14
BX462207/c 1201 bp mRNA linear EST 22-MAY-2003
LOCUS BX462207 Homo sapiens B CELLS (RAMOS CELL LINE) Homo sapiens cDNA
DEFINITION clone CSODG004YB03 5-PRIME, mRNA sequence.
ACCESSION BX462207
VERSION BX462207
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 1201)
Li,W.B., Gruber,C., Jessee,J. and Polayes,D.
Full-length cDNA libraries and normalization
Unpublished (2001)
Contact: Genoscope
Genoscope - Centre National de Sequencage
BP 191 91006 EVRY cedex - France
Email: seqref@genoscope.cns.fr, Web : www.genoscope.cns.fr
Library was constructed by Life Technologies, a division of
Invitrogen. This sequence belongs to sequence cluster 24.r For more
information about this cluster, see http://www.genoscope.cns.fr/
cgi-bin/cluster.cgi?seq=CSODG004CA02QP1&cluster=24.r. Contact :
Feng Liang Email : fliang@lifetech.com URL :

http://fulllength.invitrogen.com/ Invitrogen Corporation 1600
Faraday Avenue Genoscope sequence ID : CSODG004CA02QP1.

FEATURES
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1. .1201
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/clone="CSODG004YB03"
/tissue_type="B CELLS (RAMOS CELL LINE)"
/clone_lib="Homo sapiens B CELLS (RAMOS CELL LINE)"
/note="Vector: pCMVSPORT 6; 1st strand cDNA was primed
with a NotI-oligo(dT) primer. Five prime end enriched,
double-strand cDNA was digested with Not I and cloned into
the Not I and EcoRV sites of the pCMVSPORT 6 vector.
Library was not normalized."

ORIGIN
Query Match 18.7%; Score 42; DB 13; Length 1201;
Best Local Similarity 41.2%; Pred. No. 58;
Matches 84; Conservative 25; Mismatches 95; Indels 0; Gaps 0;

QY 22 ATAAATTTGAATAAATTATGAGGCTCCATGATGCTATATGCTTTTACCTTCAGAA 81
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Db 729 WTAATTTTAAAWAAWAAWAAWTTWWAAAAAAWTTTITTTTITTTTITTTTITTT 670

QY 82 GAATATTAGTTTCACTCAGGTTTTTCAAGCTACGCTGCCCCAAACAAACAAA 141
: : : : : : : : : : : : : : : : : : : : : : :
Db 669 AAAAAWTTTWTAAAAATTTTWTAAAAAAWTTWAAAAAAWTTAAAAATWAAAAA 610

QY 142 ACAAACAAACAACTTTTAAAGAGTTGATGGCTACTCATTTGATCGCTCCTCTGCTGA 201
: ||| ||| ||| ||| ||| : : : : : : : : : : : : : : : :
Db 609 ANAAAAAHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHHH 550

QY 202 ATCAATTAGGAATTTTITTTTITTTTITTTTITTTTITTTTITTTTITTT 225
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Db 549 TTTTITTTTITTTTITTTTITTTTITTTTITTTTITTTTITTTTITTTTITTT 526

RESULT 15
CNS058MA/c 964 bp DNA linear GSS 01-SEP-2000
LOCUS CNS058MA Tetraodon nigroviridis genome survey sequence T3 end of clone
DEFINITION 002B22 of library A from Tetraodon nigroviridis, genomic survey
sequence.
ACCESSION AL326107
VERSION AL326107.1
KEYWORDS GSS; genome survey sequence.
SOURCE Tetraodon nigroviridis
ORGANISM Tetraodon nigroviridis
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;
Tetraodontoidea; Tetraodontidae; Tetraodon.
1
Roest Crolius,H., Jaillon,O., Dasilva,C., Bouneau,L., Fisher,C.,
Bernot,A., Fizames,C., Wincker,P., Brottier,P., Quetier,F.,
Saurin,W. and Weissenbach,J.
Estimate of human gene number provided by genome-wide analysis
using Tetraodon nigroviridis DNA sequence
Nat. Genet. 25 (2), 235-238 (2000)
20296633
MEDLINE PUBMED
10835645
REFERENCE
AUTHORS Roest Crolius,H., Jaillon,O., Dasilva,C., Ozcuf-Costaz,C.,
Fizames,C., Fischer,C., Bouneau,L., Billault,A., Quetier,F.,
Saurin,W., Bernot,A. and Weissenbach,J.
TITLE Characterization and repeat analysis of the compact genome of the
freshwater pufferfish Tetraodon nigroviridis
JOURNAL Genome Res. 10 (7), 939-949 (2000)
MEDLINE PUBMED
20359837
10899143
REFERENCE
AUTHORS 3 (bases 1 to 964)

```

AUTHORS Genoscope.  
TITLE Direct Submission  
JOURNAL Submitted (12-APR-2000) Genoscope - Centre National de Sequencage :  
BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr)  
- Web : www.genoscope.cns.fr)  
COMMENT This sequence is a single read and was generated as part of a large  
scale clone-end sequencing project of the Tetraodon nigroviridis  
genome. For more information, please take a look at  
http://www.genoscope.cns.fr/Tetraodon.  
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            1..964  
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            /clone="002B22"  
            /clone\_lib="A"  
            /note="Genoscope sequence ID : C0AA002DAll1-end : T3"

## ORIGIN

Query Match 18.6%; Score 41.8; DB 29; Length 964;  
Best Local Similarity 52.4%; Pred. No. 74;  
Matches 88; Conservative 0; Mismatches 80; Indels 0; Gaps 0;  
QY 4 TGCTAAGTTGTTTCAGGCATAAATTGCAATTAATTAAGAGGCTCCATGATATGCTATA 63  
Db |||||  
821 TGGTGATTTCGGTTGTGACCATTTTTCAGCGTACAATTATGGGATTTTCTATGTTAAA 762  
QY 64 TTGGTTTACCTTCAGAGAAATATTAGTTTCACCTCAGGTTTTCAGAGCTACGCTGCC 123  
Db |||||  
761 NATTTCTACATTAAACATTATGTTTAATGTCATGAATGATGTCNAAAATTAGTTGTAT 702  
QY 124 CCCAAAAACGAAACAAACAAAAACAAACCTTTTAAAGAGTTGATG 171  
Db |||||  
701 TTAATAAAAAAACAATAATCAAGACAAGTCTCTTTTCAGATTCTG 654

Search completed: July 27, 2004, 23:03:09  
Job time : 2781 secs



GenCore version 5.1.6  
Copyright (c) 1993 - 2004 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 27, 2004, 17:03:10 ; Search time 3655 Seconds  
(without alignments)  
2668.176 Million cell updates/sec

Title: US-09-765-231a-58  
Perfect score: 225  
Sequence: 1 tgaatgtaagtgttcagg.....attaggaattttttttttt 225

Scoring table: IDENTITY NUC  
Gapop 10.0 , Gapext 1.0

Searched: 3470272 seqs, 21671516995 residues  
Total number of hits satisfying chosen parameters: 6940544

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : GenEmbl.\*

- 1: gb.ba.\*
- 2: gb.htg.\*
- 3: gb.in.\*
- 4: gb.om.\*
- 5: gb.ov.\*
- 6: gb.pat.\*
- 7: gb.ph.\*
- 8: gb.pl.\*
- 9: gb.pr.\*
- 10: gb.ro.\*
- 11: gb.sts.\*
- 12: gb.sy.\*
- 13: gb.un.\*
- 14: gb.vi.\*
- 15: em.ba.\*
- 16: em.fun.\*
- 17: em.hum.\*
- 18: em.in.\*
- 19: em.mu.\*
- 20: em.or.\*
- 21: em.ov.\*
- 22: em.pat.\*
- 23: em.ph.\*
- 24: em.pl.\*
- 25: em.ro.\*
- 26: em.sts.\*
- 27: em.un.\*
- 28: em.vi.\*
- 29: em.htg.hum.\*
- 30: em.htg.inv.\*
- 31: em.htg.other.\*
- 32: em.htg.mus.\*
- 33: em.htg.pln.\*
- 34: em.htg.rod.\*
- 35: em.htg.mam.\*
- 36: em.htg.vrt.\*
- 37: em.sy.\*
- 38: em.htgo.hum.\*
- 39: em.htgo.mus.\*
- 40: em.htgo.other.\*
- 41: em.htgo.other.\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	225	100.0	225	6	AX202128	Sequence
C 2	225	100.0	143800	2	AP000848	Homo sapi
C 3	225	100.0	186965	9	AP000848	Homo sapi
4	175.6	78.0	172830	2	AP001320	Homo sapi
5	128.4	57.1	75002	2	AC023384	Homo sapi
C 6	47.2	21.0	85916	3	AC117080	Dictyoste
7	46.2	20.5	252248	2	AC094553	Rattus no
8	45.8	20.4	110000	2	PFMAL13_04	Continuation (5 of
C 9	45.2	20.1	9971	1	U67577_01	U67577 Methanococ
10	45.2	20.1	110000	6	AR271569_01	Continuation (2 of
11	45.2	20.1	231461	2	AC096278	AC096278 Rattus no
C 12	45	20.0	170419	9	AC146265	AC146265 Pan trogl
C 13	44.4	19.7	147727	10	AL929001	AL929001 Mouse DNA
C 14	44	19.6	148750	2	AC104893	AC104893 Mus muscu
C 15	43.8	19.5	200110	10	AC117237	AC117237 Mus muscu
C 16	43	19.1	182163	2	BX000690	BX000690 Danio rer
C 17	42.6	18.9	127902	8	AP005406	AP005406 Oryza sat
C 18	42.4	18.8	199551	2	AC006281	AC006281 Plasmodiu
C 19	42.4	18.8	251551	3	AE014844	AE014844 Plasmodiu
C 20	42	18.7	147760	9	AC011846	AC011846 Homo sapi
C 21	42	18.7	151802	3	AC114263	AC114263 Dictyoste
C 22	41.8	18.6	156140	9	AC067745	AC067745 Homo sapi
C 23	41.8	18.6	215467	2	AC013420	AC013420 Homo sapi
C 24	41.8	18.6	215734	2	AC073828	AC073828 Mus muscu
C 25	41.6	18.5	171050	9	AC112232	AC112232 Homo sapi
C 26	41.6	18.5	181864	2	AC027460	AC027460 Homo sapi
C 27	41.6	18.5	194127	2	BX571853	BX571853 Danio rer
C 28	41.6	18.5	257650	2	BX323874	BX323874 Danio rer
C 29	41.4	18.4	110000	2	PFMAL7P1_03	Continuation (4 of
C 30	41.2	18.3	138564	10	AL645950	AL645950 Mouse DNA
C 31	41	18.2	5518	6	AX323692	AX323692 Sequence
C 32	41	18.2	169546	2	AC004157	AC004157 Plasmodiu
C 33	40.8	18.1	130355	8	AP003412	AP003412 Oryza sat
C 34	40.8	18.1	190561	2	AC118246	AC118246 Mus muscu
C 35	40.8	18.1	192752	2	AC118028	AC118028 Mus muscu
C 36	40.8	18.1	228071	10	AC115750	AC115750 Mus muscu
C 37	40.6	18.0	158764	2	AC132083	AC132083 Mus muscu
C 38	40.6	18.0	160429	2	AC136950	AC136950 Homo sapi
C 39	40.6	18.0	203241	9	AC008282	AC008282 Homo sapi
C 40	40.4	18.0	12300	10	MMU84903	U84903 Mus musculu
C 41	40.4	18.0	101491	10	AP003183	AP003183 Mus muscu
C 42	40.4	18.0	158123	10	AC134832	AC134832 Mus muscu
C 43	40.4	18.0	197307	2	AC130714	AC130714 Mus muscu
C 44	40.4	18.0	198600	2	AC133496	AC133496 Mus muscu
45	40.4	18.0	199371	2	AC147220	AC147220 Mus muscu

ALIGNMENTS

RESULT 1  
AX202128  
LOCUS AX202128 225 bp DNA linear PAT 30-AUG-2001  
DEFINITION Sequence 58 from Patent WO0153531.  
ACCESSION AX202128  
VERSION AX202128.1 GI:15391919  
KEYWORDS Homo sapiens (human)  
SOURCE Homo sapiens  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
REFERENCE 1  
Phippard,D., Vasanthakumari,G., Dotsen,S. and Ma,X.J.  
Osteoarthritis tissue derived nucleic acids, polypeptides, vectors,  
and cells

```
JOURNAL Patent: WO 0153531-A 58 26-JUL-2001;
Pharmacia Corporation (US)
FEATURES Location/Qualifiers
Source 1..225
/organism="Homo sapiens"
/mol_type="unassigned DNA"
/db_xref="taxon:9606"
ORIGIN
Query Match 100.0%; Score 225; DB 6; Length 225;
Best local Similarity 100.0%; Pred. No. 1.8e-40;
Matches 225; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TGATGGTAAGTGTGTTTCAGGCATAAAATTTGAATAAATATGAGGCTCCATGATGCT 60
Db 1 TGATGGTAAGTGTGTTTCAGGCATAAAATTTGAATAAATATGAGGCTCCATGATGCT 60
QY 61 ATATTGGTTTACCTTCAGAGAAATATTAGTTTCACCTCAGTGTTCACAAAGCTACGCTG 120
Db 61 ATATTGGTTTACCTTCAGAGAAATATTAGTTTCACCTCAGTGTTCACAAAGCTACGCTG 120
QY 121 TCCCCAAAACGAAACAAACAAACAAACAAACAAACAAACAAACAAACAAACAAACAAAC 180
Db 121 TCCCCAAAACGAAACAAACAAACAAACAAACAAACAAACAAACAAACAAACAAACAAAC 180
QY 181 TTGATCTGCCCTCCTCTGCTGAATCAATAGGAATTTTTCATTTTTCATTTTTCATTTT 225
Db 181 TTGATCTGCCCTCCTCTGCTGAATCAATAGGAATTTTTCATTTTTCATTTTTCATTTT 225
RESULT 2
AC011875/c 143800 bp DNA linear HTG 12-MAR-2000
LOCUS Homo sapiens clone RP11-16K5, WORKING DRAFT SEQUENCE, 26 unordered
DEFINITION pieces.
ACCESSION AC011875
VERSION AC011875.3 GI:7107950
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 143800)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,
Baldwin,J., Barna,N., Beckerly,R., Boguslavsky,L., Bouckgalter,B.,
Brown,A., Castle,A., Colangelo,M., Collins,S., Collymore,A.,
Cooke,P., DeArelano,K., Dewar,K., Domino,M., Donelan,L., Doyle,M.,
Ferreira,P., FitzHugh,W., Forrest,C., Funke,R., Gage,D.,
Gallagan,J., Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L.,
Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J.,
Lehocky,J., Lieu,C., Locke,K., Macdonald,P., Marquis,N.,
McEwan,P., McGurk,A., McKernan,K., McLaughlin,J., Meldrim,J.,
Morrow,J., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,
Peterson,K., Pollara,V., Rilev,R., Roy,A., Santos,R., Severy,P.,
Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,
Tsefaye,S., Tirrell,A., Vassiliev,H., Vo,A., Wheeler,J., Wu,X.,
Wyman,D., Ye,W.J., Zimmer,A. and Zody,M.
Direct Submission
Submitted (15-OCT-1999) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Feb 28, 2000 this sequence version replaced gi:6453961.
All repeats were identified using RepeatMasker:
Smith, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
```

```
----- Project Information
Center project name: L3566
Center clone name: 16_K_5
----- Summary Statistics
Sequencing vector: M13, M7815; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 9387 bases at least Q40
Consensus quality: 115701 bases at least Q30
Consensus quality: 130381 bases at least Q20
Insert size: 141000; agarose-fp
Insert size: 141300; sum-of-contigs
Quality coverage: 3.7 in Q20 bases; agarose-fp
Quality coverage: 3.7 in Q20 bases; sum-of-contigs
-----
* NOTE: This is a 'working draft' sequence. It currently
* consists of 26 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 1029: contig of 1029 bp in length
* 1030 1129: gap of 100 bp
* 1130 2964: contig of 1835 bp in length
* 2965 3064: gap of 100 bp
* 3065 4253: contig of 1189 bp in length
* 4254 4353: gap of 100 bp
* 4354 5678: contig of 1325 bp in length
* 5679 5779: gap of 100 bp
* 5779 6855: contig of 1077 bp in length
* 6856 6955: gap of 100 bp
* 6956 8847: contig of 1892 bp in length
* 8848 8947: gap of 100 bp
* 8948 10945: contig of 1998 bp in length
* 10946 11045: gap of 100 bp
* 11046 13001: contig of 1956 bp in length
* 13002 13101: gap of 100 bp
* 13102 15430: contig of 2329 bp in length
* 15431 15530: gap of 100 bp
* 15531 17919: contig of 2389 bp in length
* 17920 18019: gap of 100 bp
* 18020 20135: contig of 2116 bp in length
* 20136 20235: gap of 100 bp
* 20236 23750: contig of 3515 bp in length
* 23751 23850: gap of 100 bp
* 23851 26794: contig of 2944 bp in length
* 26795 26894: gap of 100 bp
* 26895 29482: contig of 2588 bp in length
* 29483 29582: gap of 100 bp
* 29583 33174: contig of 3592 bp in length
* 33175 33274: gap of 100 bp
* 33275 35185: contig of 1911 bp in length
* 35186 35285: gap of 100 bp
* 35286 39745: contig of 4460 bp in length
* 39746 39845: gap of 100 bp
* 39846 44222: contig of 4377 bp in length
* 44223 48990: contig of 4668 bp in length
* 48991 49090: gap of 100 bp
* 49091 57790: contig of 8700 bp in length
* 57791 57890: gap of 100 bp
* 57891 66822: contig of 8932 bp in length
* 66823 66922: gap of 100 bp
* 66923 76709: contig of 9787 bp in length
* 76710 76809: gap of 100 bp
* 76810 92865: contig of 16056 bp in length
* 92866 92966: gap of 100 bp
* 92967 106278: contig of 13313 bp in length
* 106279 106378: gap of 100 bp
* 106379 123041: contig of 16663 bp in length
* 123042 123141: gap of 100 bp
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\* 165955 165954: gap of 100 bp  
 \* 165955 167495: contig of 1541 bp in length  
 \* 167496 167595: gap of 100 bp  
 \* 167596 168771: contig of 1176 bp in length  
 \* 168772 168871: gap of 100 bp  
 \* 168872 170322: contig of 1451 bp in length  
 \* 170323 170422: gap of 100 bp  
 \* 170423 171562: contig of 1140 bp in length  
 \* 171563 171562: gap of 100 bp  
 \* 171663 172830: contig of 1168 bp in length.

## FEATURES

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 /map="11q14"  
 /clone="RP11-79904"

misc\_feature  
 1..15732  
 /note="assembly\_fragment"  
 15833..34101  
 /note="assembly\_fragment"  
 34202..47479  
 /note="assembly\_fragment"  
 47580..57756  
 /note="assembly\_fragment"  
 57857..69426  
 /note="assembly\_fragment"  
 69527..80185  
 /note="assembly\_fragment"  
 80286..88989  
 /note="assembly\_fragment"  
 89090..99362  
 /note="assembly\_fragment"  
 99463..103433  
 /note="assembly\_fragment clone\_end:SP6 vector\_side:left"  
 103534..109064  
 /note="assembly\_fragment"  
 109165..113855  
 /note="assembly\_fragment"  
 113956..119140  
 /note="assembly\_fragment"  
 119241..124661  
 /note="assembly\_fragment"  
 124762..126822  
 /note="assembly\_fragment clone\_end:T7 vector\_side:right"  
 126923..131337  
 /note="assembly\_fragment"  
 131438..135078  
 /note="assembly\_fragment"  
 135179..138134  
 /note="assembly\_fragment"  
 138235..142123  
 /note="assembly\_fragment"  
 142224..144719

Query Match 78.0%; Score 175.6; DB 2; Length 172830;  
 Best Local Similarity 88.9%; Pred. No. 6,7e-30;  
 Matches 201; Conservative 0; Mismatches 24; Indels 1; Gaps 1;

QY 1 TGATGGTAACTGTTTCAGGCATTAATTTCAATAAATATGAGGCTCCATGATAGCT 60  
 Db 28909 TGATGGTAACTGTTTCAGGCATTAATTTCAATAAATATGAGGCTCCATGATAGCT 28968  
 QY 61 ATATTGTTTACCTTCAGAGATATTTAGTTTCACTCAGGTTTTTCAAGCT-ACGCT 119  
 Db 28969 ATATTGTTTACCTTCAGAGATATTTAGTTTCACTCAGGTTTTTCAAGCTAACCT 29028  
 QY 120 GTCCCCCAAAACGAAACAAAACAAACCTTTTAAAGAGTTGATGCGTACTCA 179  
 Db 29029 GTCCCCCAAAACGAAACAAAACCTTTTAAAGAGTTGATGCGTAAATCA 29088  
 QY 180 TTGATCTGCTCTCTGCTGAATCAATTAGGAATTTTTTTTTT 225

Db 29089 TTGATTTGCTCTGCTGTCGATCACATTAGGGATTTTCTTTT 29134

## RESULT 5

AC023384

LOCUS

DEFINITION

AC023384

VERSION

AC023384.2

KEYWORDS

HTG; HTGS\_PHASE0.

SOURCE

Homo sapiens

ORGANISM

Homo sapiens

REFERENCE

1 (bases 1 to 75002)

AUTHORS

Birren,B., Linton,L., Nusbaum,C. and Lander,E.

TITLE

Homo sapiens chromosome 11, clone RP11-589112

JOURNAL

Unpublished

REFERENCE

2 (bases 1 to 75002)

AUTHORS

Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,

Anderson,S., Baldwin,J., Barna,N., Beda,F., Boguslavskiy,L.,

Boukhgalter,B., Brown,A., Burkett,G., Campopiano,A., Castle,A.,

Choepe,Y., Collangelo,M., Collins,S., Collamore,A., Cooke,P.,

DeArellano,K., Dewar,K., Dodge,S., Domino,M., Doyle,M.,

Fenestor,J., Ferreira,P., FitzHugh,W., Forrest,C., Gage,D.,

Galagan,J., Gardyna,S., Ginde,S., Goyette,M., Graham,L.,

Grand-Pierre,N., Grant,G., Hagos,B., Heaford,A., Horton,L.,

Howland,J.C., Iliev,I., Johnson,R., Jones,C., Kann,L., Karatas,A.,

Klein,J., Landers,T., Largocque,K., Lehoczy,J., Levine,R.,

Lieu,C., Liu,G., Locke,K., Macdonald,P., Marquis,N., McCarthy,M.,

McEwan,P., McGurk,A., McKernan,K., McPheeters,R., Meldrim,J.,

Meneus,L., Mihova,T., Miranda,C., Menga,V., Morrow,J., Naylor,J.,

Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Olivari,T.M.,

Peterson,K., Pierre,N., Pisani,C., Pollara,V., Raymond,C.,

Riley,R., Rogov,P., Rothman,D., Roy,A., Santos,R., Schauer,S.,

Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N.,

Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J., Tirrell,A.,

Travers,M., Trigilio,J., Vassiliev,H., Viel,R., Vo,A., Wilson,B.,

Wu,X., Wyman,D., Ye,W.J., Young,G., Zainoun,J., Zimmer,A. and

Zody,M.

Direct Submission

Submitted (14-FEB-2000) Whitehead Institute/MIT Center for Genome

Research, 320 Charles Street, Cambridge, MA 02141, USA

On Jul 13, 2000 this sequence version replaced gi:6970532.

All repeats were identified using RepeatMasker:

Smit, A.F.A. &amp; Green, P. (1996-1997)

http://ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBR

Web site: http://www-seq.wi.mit.edu

Contact: sequence\_submissions@genome.wi.mit.edu

----- Project Information

Center project name: L5114

-----

\* NOTE: This record contains 88 individual

\* sequencing reads that have not been assembled into

\* contigs. Runs of N are used to separate the reads

\* and the order in which they appear is completely

\* arbitrary. Low-pass sequence sampling is useful for

\* identifying clones that may be gene-rich and allows

\* overlap relationships among clones to be deduced.

\* However, it should not be assumed that this clone

\* will be sequenced to completion. In the event that

\* the record is updated, the accession number will

\* be preserved.

\* 1 724: contig of 724 bp in length

\* 725 824: gap of 100 bp

\* 825 1573: contig of 749 bp in length

\* 1574 1673: gap of 100 bp



```
QY      204 CAATTAGGAATTTTTTTTTT 225
LOCUS   AC117080/c
DEFINITION Dictyostelium discoideum chromosome 2 map complement (821514-735598)
strain AX4, complete sequence.
ACCESSION AC117080 AC114260
VERSION   AC117080.2 GI:28850277
KEYWORDS  HTG.
SOURCE    Dictyostelium discoideum
ORGANISM  Eukaryota; Mycetozoa; Dictyosteliida; Dictyostelium.
REFERENCE 1 (bases 1 to 85916)
AUTHORS   Gloeckner,G., Eichinger,L., Szafranski,K., Rachebat,J., Dear,P.,
           Lehmann,R., Baumgart,C., Parra,G., April,J.F., Guigo,R., Kumpf,K.,
           Tungal,B., Cox,E., Quail,M.A., Platzer,M., Rosenthal,A. and
           Noegel,A.A.
TITLE      Sequence and analysis of chromosome 2 of Dictyostelium discoideum
JOURNAL   Nature 418 (6893), 79-85 (2002)
MEDLINE   22092622
PUBMED    12097910
REMARK    The Dictyostelium Genome Sequencing Consortium
REFERENCE 2 (bases 1 to 85916)
AUTHORS   Baumgart,C.
TITLE      Direct Submission
JOURNAL   Submitted (06-APR-2002) Genome Analysis, Institute of Molecular
           Biotechnology, Beutenbergstr. 11, Jena 07745, Germany
REFERENCE 3 (bases 1 to 85916)
AUTHORS   Baumgart,C.
TITLE      Direct Submission
JOURNAL   Submitted (05-MAR-2003) Genome Analysis, Institute of Molecular
           Biotechnology, Beutenbergstr. 11, Jena 07745, Germany
REFERENCE 4 (bases 1 to 85916)
AUTHORS   Baumgart,C.
TITLE      Direct Submission
JOURNAL   Submitted (12-MAR-2003) Genome Analysis, Institute of Molecular
           Biotechnology, Beutenbergstr. 11, Jena 07745, Germany
COMMENT   On or before Mar 5, 2003 this sequence version replaced
           gi:13424360, gi:20066270.
CDS predictions from GeneID do not necessarily reflect true genes.
Further information is available from IMB Jena, Department of
Genome Analysis
(http://genome.imb-jena.de/dictyostelium/)
and the University Cologne, Institute for Biochemistry I
(http://www.uni-koeln.de/dictyostelium/project.shtml)
Funding
Agency : Deutsche Forschungsgemeinschaft (DFG).
           Location/Qualifiers
           1..85916
              /organism="Dictyostelium discoideum"
              /mol_type="genomic DNA"
              /strain="AX4"
              /db_xref="taxon:44689"
              /map="complement (821514-735598)"
              /chromosome="2"
              join(9..188,279..1337)
              /note="GeneID exon scores (in order of location ranges):
              9,57, 89,77 - GSCJ_ID dd_00785"
              /codon_start=1
              /product="hypothetical protein"
              /protein_id="AA053082.1"
              /db_xref="GI:28850278"
              /translation="MEDNNSNNENKQNELSTPSTIKERHVLVIMQHLGHTSLDPKT
              IRNHLKQKLDNCIFTSANSNSHFLATHDGIDKIGERLFNEKELYEQYDHPKISM
              IGHSGLGLITRYAIGLLYDDGFFKCKPDQFI SLSPHCGSRFPSTTFINKVAHIFVD
              NFLVSGKQILHDEITPDNIKTFFPSTSPPPNEKLSKSTIVNSVKNETDLSPLA
              EAKPSTYKEVGNEKMLIETKEENEIITNDQVPMPLAVRLSEGIFFENGLNSFRK
              TLYSNINDVQVNFCTSDISAKNPYTLGKLMKFSKRYHRIIEESILDIDPNLLEQOS
              PPPDKPFDEKLDDEYFTHDTHHHLKRLKLNQLHFVRYHYFMKMLSHNTNIIVKR
```

```
EWINSEGFIIIEHLVSHPEG"
join(2619..2689,2806..2917,3021..3235,3389..3938)
/note="GeneID exon scores (in order of location ranges):
4,00, 11,19, 30,82, 62,55 - GSCJ_ID dd_00787"
/codon_start=1
/product="similar to Mus musculus (Mouse). 26S proteasome
regulatory subunit 55A (Rpn10) (Multiubiquitin chain
binding protein)"
/protein_id="AA053083.1"
/db_xref="GI:28850279"
/translation="MYTASKTQSNPESAVSIMSMACKPEVLVTLTQLSKLSKLGAE
IKINGKIDSTTMOIALALRHQNNHQPRIAFVGSPLKETKEELIQLAKEKNG
VAVDLINFGEVNSDKLEAFINDVNNDESHLLTPPGPHILSDILQSPIVSGGS
QFGSEFINATDPTDLAMALKISLEEKQERERKAREBANGSTLTPTTTPATES
NMDVNFDDPELAALSMATDKMEVQSSTNTDSQPPQQQQPPTDDTSTSEAFKQD
DFNLSTNLSPGVDPNRIKNALENLSKDEDKDKNEKK"
complement(join(4186..5395,5501..5689,5854..6303,
6445..7055))
/note="GeneID exon scores (in order of location ranges):
94,07, 12,83, 33,44, 41,83 - GSCJ_ID dd_00788"
/codon_start=1
/product="similar to Mus musculus (Mouse). 12 days embryo
male wolffian duct includes surrounding region cDNA, RIKEN
full-length enriched library, clone:6720456H20
product:similar to KAI1A4653 PROTEIN (Fragment)"
/protein_id="AAM45299.2"
/db_xref="GI:28850280"
/translation="MNSPTLSNTRNRKNNNSNSNSNNNNNNNNNNNNNNNNNN
NNNNVNRVNRNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNN
RFGKLLKSTIACIVLLFTSCIIYSMTQPSVSGDAGELIVAHQGVNHPGYPVL
FTFLGFIFSHIIPSNLTSVANKISFMSMIGSITASTIYTLVILWNHNNHGWGLSAYM
FSPSLIMWYQIQGEVFSNMNMFVAMLFGLVWYTRVIFENERYNTAFWTSRIAYL
FTPKLNRVKYSGMNSTLRGFIKHEFREYGTLYQDGVGVSITKISIVENLI
IQGYTGLALSLIGLLNLGNIRTFKWSFGCTMII FSLFYITFENLGNIPDKP
LYRGVPLRFPMPQNVLIITMGLGIKSIQFLNRIDQDGGTATTTTISKIQKLLP
IIILLVGNQIQNYLQDQSNISFYDSVLDGLPRLNLLVGLDGLVNVPMYLIH
LCEKRPDIDILSMEIMSWEFKTSIPLQRFQVKGPNVHPYIPGYSLSKLDANI
NRPYIYIGGDFKSGSFQNDYITI SKLTSQIIPKDSHKENTFKIITFTFSPF
HIPNPKYFNDPSWHEFMEMAVSLERAETLLKEYLSQNTSEKALESLVEILEK
AIITYNDKQWSLKHGICFDHLRYRVQVNNNNNQAKNSYKQLLYFWKKYINQ
CTHEENTDQDMETIKKVIQLI"
7833..8195
/note="GeneID exon scores (in order of location ranges):
30,55 - GSCJ_ID dd_00789"
/codon_start=1
/product="hypothetical protein"
/protein_id="AA053084.1"
/db_xref="GI:28850281"
/translation="MINLKNKPVLTGLMNSVFFPNIKLVIRNSKGTENTNMKIFK
TDCSVGKTIDIKTYMKALYDVNVKNTINVQGRIKSTKGLQKRSKLSKYKTPDYK
AIITVDPSLRAQLSRGKN"
9000..9674
/note="GeneID exon scores (in order of location ranges):
54,39 - GSCJ_ID dd_02815"
/codon_start=1
/product="similar to Homo sapiens (Human). Small GTP
binding protein RAB32"
/protein_id="AAM45300.2"
/db_xref="GI:28850282"
/translation="MYSNKNKDKDKDKQNNNNNNNDDEAISLIILVSKLACGKTSI
IORYCHNNFQPKYPTIGVDFQOKVLEIMQKVLQIQLWDIAGERGHMTRVFFQAH
GAVIVFDATRSGLTFLGAKWKDDIDYCFNNENLPTLLANKCDLLTPPYTFPEDINTF
CEONRFNKYFVTSKEDTGINALEELVLKILLESYQTEQSTGFKLSDQSSTETPTT
QSKTCC"
complement(10728..12734)
/note="GeneID exon scores (in order of location ranges):
120,50 - GSCJ_ID dd_03061"
/codon_start=1
/product="similar to Plasmodium falciparum. Hypothetical
protein"
/protein_id="AA053085.1"
/db_xref="GI:28850283"
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TITLE  
JOURNAL  
REFERENCE  
AUTHORS

Direct Submission  
Unpublished  
2 (bases 1 to 252248)  
Worley,K.C.

TITLE  
JOURNAL

Submitted (15-SEP-2001) Human Genome Sequencing Center, Department  
of Molecular and Human Genetics, Baylor College of Medicine, One  
Baylor Plaza, Houston, TX 77030, USA  
3 (bases 1 to 252248)  
Rat Genome Sequencing Consortium.

REFERENCE  
AUTHORS

Direct Submission  
Submitted (09-MAY-2003) Human Genome Sequencing Center, Department  
of Molecular and Human Genetics, Baylor College of Medicine, One  
Baylor Plaza, Houston, TX 77030, USA

TITLE  
JOURNAL

## COMMENT

The sequence in this assembly is a combination of BAC based reads  
and whole genome shotgun sequencing reads assembled using Atlas  
(<http://www.hgsc.bcm.tmc.edu/projects/rat/>). Each contig described  
in the feature table below represents a scaffold in the Atlas  
assembly (a 'contig-scaffold'). Within each contig-scaffold,  
individual sequence contigs are ordered and oriented, and separated  
by sized gaps filled with Ns to the estimated size. The sequence  
may extend beyond the ends of the clone and there may be sequence  
contigs within a contig-scaffold that consist entirely of whole  
genome shotgun sequence reads. Both end sequences and whole genome  
shotgun sequence only contigs will be indicated in the feature  
table.

----- Genome Center  
Center: Baylor College of Medicine

Center code: BCM

Web site: <http://www.hgsc.bcm.tmc.edu/>

Contact: [hgsc-help@bcm.tmc.edu](mailto:hgsc-help@bcm.tmc.edu)

----- Project Information

Center project name: GAWO

Center clone name: CH230-4P5

----- Summary Statistics

Assembly program: Atlas;

Consensus quality: 225835 bases at least Q40

Consensus quality: 228048 bases at least Q30

Consensus quality: 229440 bases at least Q20

Estimated insert size: 233224; sum-of-contigs estimation

Quality coverage: 7x in Q20 bases; sum-of-contigs estimation

-----  
\* NOTE: Estimated insert size may differ from sequence length  
(see [http://www.hgsc.bcm.tmc.edu/docs/genbank\\_draft\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html)).  
\* NOTE: This is a 'working draft' sequence. It currently  
\* consists of 5 contigs. The true order of the pieces  
\* is not known and their order in this sequence record is  
\* arbitrary. Gaps between the contigs are represented as  
\* runs of N, but the exact sizes of the gaps are unknown.  
\* This record will be updated with the finished sequence  
\* as soon as it is available and the accession number will  
\* be preserved.

\* 1 246259: contig of 246259 bp in length

\* 246260 246359: gap of unknown length

\* 246360 247679: contig of 1320 bp in length

\* 247680 247779: gap of unknown length

\* 247780 249446: contig of 1667 bp in length

\* 249447 249546: gap of unknown length

\* 249547 250637: contig of 1091 bp in length

\* 250638 250737: gap of unknown length

\* 250738 252248: contig of 1511 bp in length.

## FEATURES

## source

1. .252248

/organism="Rattus norvegicus"

/mol\_type="genomic DNA"

/db\_xref="taxon:10116"

/clone="CH230-4P5"

1. .2282

/note="wgs end\_extension"

clone\_end.T7"

7986..8866

/note="clone\_boundary"

## misc\_feature

clone\_end:T7  
site:EcoRI  
end sequence:BH310954"  
240861..242054

## misc\_feature

/note="wgs contig"  
complement(240876..241519)  
/note="clone boundary"

## misc\_feature

clone\_end:Sp6  
site:EcoRI  
end sequence:BH310955"  
243149..246259  
/note="wgs end\_extension"

## ORIGIN

Query Match 20.5%; Score 46.2; DB 2; Length 252248;

Best Local Similarity 54.4%; Pred. No. 0.62; 78; Indels 0; Gaps 0;

Matches 93; Conservative 0; Mismatches 78; Indels 0; Gaps 0;

QY 24 AAAATTGAAATAAATTATGAGGCTCCATGATATGCTATATGCTTTTACCTTCAGAGA 83

Db 205591 AAAATTGCTTCAATTTCTTTGATACAAAGATTGAGGCTTTTATTTTCCCATAGAGAA 205650

QY 84 ATATTAGTTTCACTCAGGTTTTTCAAGCTACGCTGTCCCAAAAAACGAAACAAAC 143

Db 205651 TGCTTGAATTTCACTAAGAAATTTTCAAGTACATTTAGTGTCTGAAAAAAGAAAG 205710

QY 144 AAAAAAACCAACCTTTTAAAGATTGATGGCTACTCATTTGATCTGCTCTCT 194

Db 205711 AAAAGAAGAAAGAAATTTCCAGTTGATTGTTACTAATTAAGAATTTCTCTT 205761

RESULT 8

PFMAL13\_04

WPCOMMENT

Sequence split into 28 fragments LOCUS PFMAL13 Accession AL844509

Fragment Name Begin End

PFMAL13\_00 1 110000

PFMAL13\_01 100001 210000

PFMAL13\_02 200001 310000

PFMAL13\_03 300001 410000

PFMAL13\_04 400001 510000

PFMAL13\_05 500001 610000

PFMAL13\_06 600001 710000

PFMAL13\_07 700001 810000

PFMAL13\_08 800001 910000

PFMAL13\_09 900001 1010000

PFMAL13\_10 1000001 1110000

PFMAL13\_11 1100001 1210000

PFMAL13\_12 1200001 1310000

PFMAL13\_13 1300001 1410000

PFMAL13\_14 1400001 1510000

PFMAL13\_15 1500001 1610000

PFMAL13\_16 1600001 1710000

PFMAL13\_17 1700001 1810000

PFMAL13\_18 1800001 1910000

PFMAL13\_19 1900001 2010000

PFMAL13\_20 2000001 2110000

PFMAL13\_21 2100001 2210000

PFMAL13\_22 2200001 2310000

PFMAL13\_23 2300001 2410000

PFMAL13\_24 2400001 2510000

PFMAL13\_25 2500001 2610000

PFMAL13\_26 2600001 2710000

PFMAL13\_27 2700001 2732359

Continuation (5 of 28) of PFMAL13 from base 400001 (AL844509 Plasmodium falciparum 3D7 c

Query Match 20.4%; Score 45.8; DB 2; Length 110000;

Best Local Similarity 51.2%; Pred. No. 0.87;

Matches 107; Conservative 0; Mismatches 102; Indels 0; Gaps 0;

QY 17 CAGGCATATAAATTGAAATAAATTATGAGGCTCCATGATATGCTATATGCTTTTACCTT 76

Db 28080 CAGCTATAAAATTTACACGTCCTATTATATATACACATAAAATATATATATATATATA 28139

Qy 77 CAGAAGAAATATTAGTTTCACTCAGGTTTTCCTCAAGCTACGCTGCCCCAAAACCAA 136

Db 28140 TATATATATATTATATATATAAAATATTTTGAACACTAGCGAAAAAATAAAAAA 28199

Qy 137 AAAAAACAAAAAACAACCTTTTAAAGATGTCAGGCTACTCAITTTGATCTGCCTCTCT 196

Db 28200 AAAAAAATAAATAAGCTTTTAAATAATATTTTCCCTGCTGTTTCAATGATTTTAA 28259

Qy 197 GCTGATCAATAGGAATTTTTTTTTTTT 225

Db 28260 AATAAGAAATATTTTTTTTTTTTTTTTTTTT 28288

RESULT 9

U67577/c

LOCUS U67577 9971 bp DNA linear BCT 28-JAN-1998

DEFINITION Methanococcus jannaschii section 119 of 150 of the complete genome.

ACCESSION U67577 L77117

VERSION U67577.1 GI:2826400

KEYWORDS

SOURCE Methanocaldococcus jannaschii

ORGANISM Methanocaldococcus jannaschii

Archaea; Euryarchaeota; Methanococci; Methanococcales; Methanocaldococaceae; Methanocaldococcus.

REFERENCE 1 (bases 1 to 9971)

AUTHORS Bult, C.J., White, O., Olsen, G.J., Zhou, L., Fleischmann, R.D., Sutton, G.G., Blake, J.A., Fitzgerald, L.M., Clayton, R.A., Gocayne, J.D., Kerlavage, A.R., Dougherty, B.A., Tomb, J., Adams, M.D., Reich, C.I., Overbeek, R., Kirkness, E.F., Weinstock, K.G., Merrick, J.M., Glodek, A., Scott, J.D., Geoghagen, N.S., Weidman, J.F., Fuhrmann, J.L., Nguyen, D.T., Utterback, T., Kelley, J.M., Peterson, J.D., Sadow, P.W., Hanna, M.C., Cotton, M.D., Hurst, M.A., Roberts, K.M., Kaine, B.B., Borodovsky, M., Klenk, H.P., Fraser, C.M., Smith, H.O., Woese, C.R. and Venter, J.C.

TITLE Complete genome sequence of the methanogenic archaeon, Methanococcus jannaschii

JOURNAL Science 273 (5278), 1058-1073 (1996)

MEDLINE 96337999

PUBMED 8688087

REFERENCE 2 (bases 1 to 9971)

AUTHORS Bult, C.J., White, O., Olsen, G.J., Zhou, L., Fleischmann, R.D., Sutton, G.G., Blake, J.A., Fitzgerald, L.M., Clayton, R.A., Gocayne, J.D., Kerlavage, A.R., Dougherty, B.A., Tomb, J., F., Adams, M.D., Reich, C.I., Overbeek, R., Kirkness, E.F., Weinstock, K.G., Merrick, J.M., Glodek, A., Scott, J.D., Geoghagen, N.S., Weidman, J.F., Fuhrmann, J.L., Nguyen, D.T., Utterback, T., Kelley, J.M., Peterson, J.D., Sadow, P.W., Hanna, M.C., Cotton, M.D., Hurst, M.A., Roberts, K.M., Kaine, B.B., Borodovsky, M., Klenk, H.P., Fraser, C.M., Smith, H.O., Woese, C.R. and Venter, J.C.

Direct Submission

Submitted (27-AUG-1996) The Institute for Genomic Research, 9712 Medical Center Dr, Rockville, MD 20850, USA

COMMENT On Jan 30, 1998 this sequence version replaced gi:1592013.

FEATURES

source

1. .9971

/organism="Methanocaldococcus jannaschii"

/mol\_type="genomic DNA"

/db\_xref="taxon:2190"

87. .554

/gene="MJ1369"

87. .554

/note="MJ1369"

/note="hypothetical protein; identified by GeneMark; putative"

/codon\_start=1

/transl\_table=11

/product="M. jannaschii predicted coding region MJ1369"

/protein\_id="AAB99386.1"

/db\_xref="GI:1592014"

/translation="MRDVMYTWLFFSSNNIRNIEICYNHMLGFWDFNAGEKOKNWR

gene

CDS

SFSVLIIFSKEAVIKRFIKIQDYIDCYGLGELEHDFNEILKAFQKFGWISIK"

584. .1570

/gene="MJ1370"

584. .1570

/gene="MJ1370"

/note="hypothetical protein; identified by GeneMark; putative"

/codon\_start=1

/transl\_table=11

/product="M. jannaschii predicted coding region MJ1370"

/protein\_id="AAB99387.1"

/db\_xref="GI:1592015"

/translation="MITVGDHGTSGITTCIKNDKKIIFKLKTELKESYLEBELEK HISLEDIDLALTYMGDGINKILPIEKKNRGVLSLEGAGEKVGKGGTKVYDEIKESG LPVAVIPGLHGTIECLDERFINKLYSHIASPEKVSIAIYAYKLFENFPVLSIDISNTV TLLIKDGKIFGGFDACIGAGMLHPIDLEIMIRIDAGKITANEAFSKAGAVKIANKLY KGNTKEEIRINNYDENDECLAVDSLIVSMEINSLPLDLLDKRRRVLAGISGTL RNPTIDIPKRIEFVEAKIFVLYGSGAIGGALIAEDILKGRDILGIEVEFK"

1567. .2301

/gene="MJ1371"

1567. .2301

/gene="MJ1371"

/note="similar to GP:2437836 percent identity: 39.09; identified by sequence similarity; putative"

/codon\_start=1

/transl\_table=11

/product="methyltransferase"

/protein\_id="AAB99379.1"

/db\_xref="GI:1592016"

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2285. .3127

/gene="MJ1372"

2285. .3127

/gene="MJ1372"

/note="similar to GB:L42023 SP:P44938 PID:1006041 PID:1221025 PID:1205167 percent identity: 47.86; identified by sequence similarity; putative"

/codon\_start=1

/transl\_table=11

/product="conserved hypothetical protein"

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/db\_xref="GI:1592017"

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3124. .3612

/gene="MJ1373"

3124. .3612

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/note="hypothetical protein; identified by GeneMark; putative"

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3680. .4480

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3680. .4480

/gene="MJ1374"

/note="similar to GB:AB000666 percent identity: 42.73; identified by sequence similarity; putative"

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CDS

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3124. .3612

/gene="MJ1373"

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/gene="MJ1374"

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/gene="MJ1374"

3680. .4480

/gene="MJ1374"

/note="similar to GB:AB000666 percent identity: 42.73; identified by sequence similarity; putative"



## ORGANISM

Rattus norvegicus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;  
Rattus.

REFERENCE  
AUTHORS

1 (bases 1 to 231461)  
Murny,D.Marie., Metzker,M.Lee., Abramzon,S., Adams,C., Alder,J.,  
Allen,C., Allen,H., Albrooks,S., Amin,A., Anguiano,D.,  
Anyalebechi,V., Aoyagi,A., Ayodeji,M., Baca,E., Baden,H.,  
Baldwin,D., Bandaranaike,D., Barber,M., Barnstead,M., Benahmed,P.,  
Biswal,K., Blair,J., Burch,P., Blankenburg,K., Blyth,P., Brown,M.,  
Bryant,N., Buhay,C., Burch,P., Burrell,K., Calderon,E.,  
Cardenas,V., Carter,K., Cavazos,I., Ceasar,H., Center,A.,  
Chacko,J., Chavez,D., Chen,G., Chen,R., Chen,Y., Chen,Z., Chu,J.,  
Cleveland,C., Cockrell,R., Cox,C., Coyle,M., Cree,A., D'Souza,L.,  
Davila,M.L., Davis,C., Davy-Carroll,L., De Anda,C., Dederich,D.,  
Delgado,O., Denson,S., Deramo,C., Ding,Y., Dinh,H., Divya,K.,  
Draper,H., Dugan-Rocha,S., Dunn,A., Durbin,K., Duval,B., Eaves,K.,  
Egan,A., Escotto,M., Eugene,C., Evans,C.A., Falls,T., Fan,G.,  
Fernandez,S., Finley,M., Flagg,N., Forbes,L., Foster,M., Foster,P.,  
Fraser,C.M., Gabisi,A., Ganta,R., Garcia,A., Garner,T., Garza,M.,  
Gebregorgis,E., Geer,K., Gill,R., Grady,M., Guerra,W., Guevara,W.,  
Gunaratne,P., Haaland,W., Hamil,C., Hamilton,C., Hamilton,K.,  
Harvey,Y., Havlak,P., Hawes,A., Henderson,N., Hernandez,J.,  
Hernandez,R., Hines,S., Hladun,S.L., Hodgson,A., Hogues,M.,  
Hollins,B., Howells,S., Huliyk,S., Hume,J., Idlebird,D., Jackson,A.,  
Jackson,L., Jacob,L., Jiang,H., Johnson,B., Johnson,R., Jolivet,A.,  
Karpachy,S., Kelly,S., Kelly,S., Khan,Z., King,L., Kovar,C.,  
Kovis,C., Kraft,C.L., Lebow,H., Levan,J., Lewis,L., Li,Z., Liu,J.,  
Liu,J., Liu,W., Liu,Y., London,P., Longacre,S., Lopez,J.,  
Lorensuhewa,B., Louised,H., Lozado,R.J., Lu,X., Ma,J.,  
Maheshwari,M., Mahindartine,M., Mahmoud,M., Malloy,K., Mangum,A.,  
Mangum,B., Mapua,P., Martin,K., Martin,R., Martinez,E.,  
Mahoney,S., McLeod,M.P., McNeill,T.Z., Meenen,E.,  
Milosavljevic,A., Miner,G., Minja,E., Montemayor,J., Moore,S.,  
Morgan,M., Morris,K., Morris,S., Munidasa,M., Murphy,M., Nair,L.,  
Nankervis,C., Neal,D., Newton,N., Nguyen,N., Norris,S.,  
Nwackemeleh,O., Okwuonu,G., Olarnpunsagoon,A., Pal,S., Parks,K.,  
Pasternak,S., Paul,H., Perez,A., Perez,L., Pfamkoch,C.,  
Plopper,F., Poidexter,A., Popovic,D., Primus,E., Pu,L.-L.,  
Pruzo,M., Quiroz,J., Rachlin,E., Reeves,K., Regier,M.A., Reigh,R.,  
Reilly,B., Reilly,M., Ren,Y., Reuter,M., Richards,S., Riggs,F.,  
Rivers,C., Rodkey,T., Rojas,A., Rose,M., Rose,R., Ruiz,S.J.,  
Sanders,W., Savery,G., Scherer,S., Scott,G., Shatsman,S., Shen,H.,  
Shetty,J., Shvartsbeyn,A., Sisson,I., Sitter,C.D., Smajs,D.,  
Sneed,A., Sodergren,E., Song,X.-Z., Sorelle,R., Sosa,J.,  
Steimle,M., Strong,R., Sutton,A., Svatek,A., Tabor,P., Taylor,C.,  
Taylor,T., Thomas,N., Thomas,S., Tingey,A., Trejos,Z., Usmani,K.,  
Valas,R., Vera,V., Villasana,D., Waldron,L., Walker,B., Wang,J.,  
Wang,Q., Wang,S., Warren,J., Warren,R., Wei,X., White,F.,  
Williams,G., Willson,R., Wleczyk,R., Wooden,H., Worley,K.,  
Wright,D., Wright,R., Wu,J., Yakub,S., Yen,J., Yoon,L., Yoon,V.,  
Yu,F., Zhang,J., Zhou,J., Zhou,X., Zhao,S., Dunn,D., von  
Niederhausen,A., Weiss,R., Smith,D.R., Holt,R.A., Smith,H.O.,  
Weinstock,G. and Gibbs,R.A.

## TITLE

Direct Submission

## JOURNAL

2 (bases 1 to 231461)  
Unpublished

## REFERENCE

Worley,K.C.

## AUTHORS

Worley,K.C.

## JOURNAL

Submitted (17-SEP-2001) Human Genome Sequencing Center, Department  
of Molecular and Human Genetics, Baylor College of Medicine, One  
Baylor Plaza, Houston, TX 77030, USA

## REFERENCE

3 (bases 1 to 231461)

## AUTHORS

Rat Genome Sequencing Consortium.

## TITLE

Direct Submission

## JOURNAL

Submitted (10-MAY-2003) Human Genome Sequencing Center, Department  
of Molecular and Human Genetics, Baylor College of Medicine, One  
Baylor Plaza, Houston, TX 77030, USA

## COMMENT

The sequence in this assembly is a combination of BAC based reads  
and whole genome shotgun sequencing reads assembled using Atlas  
(<http://www.hgsc.bcm.tmc.edu/projects/rat/>). Each contig described  
in the feature table below represents a scaffold in the Atlas

assembly (a 'contig-scaffold'). Within each contig-scaffold,  
individual sequence contigs are ordered and oriented, and separated  
by sized gaps filled with 'Ns' to the estimated size. The sequence  
may extend beyond the ends of the clone and there may be sequence  
contigs within a contig-scaffold that consist entirely of whole  
genome shotgun sequence reads. Both end sequences and whole genome  
shotgun sequence only contigs will be indicated in the feature  
table.

----- Genome Center  
Center: Baylor College of Medicine  
Center code: BCM  
Web site: <http://www.hgsc.bcm.tmc.edu/>  
Contact: [hgsc-help@bcm.tmc.edu](mailto:hgsc-help@bcm.tmc.edu)  
----- Project Information  
Center project name: GEVR  
Center clone name: CH230-11L24  
----- Summary Statistics  
Assembly program: Atlas 3.0;  
Consensus quality: 225324 bases at least Q40  
Consensus quality: 227015 bases at least Q30  
Consensus quality: 227940 bases at least Q20  
Estimated insert size: 232676; sum-of-contigs estimation  
Quality coverage: 8x in Q20 bases; sum-of-contigs estimation  
-----  
\* NOTE: Estimated insert size may differ from sequence length  
(see [http://www.hgsc.bcm.tmc.edu/docs/Genbank\\_draft\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html)).  
\* NOTE: This is a 'working draft' sequence. It currently  
\* consists of 1 contigs. Gaps between the contigs  
\* are represented as runs of N. The order of the pieces  
\* is believed to be correct as given, however the sizes  
\* of the gaps between them are based on estimates that have  
\* provided by the submitter.  
\* This sequence will be replaced  
\* by the finished sequence as soon as it is available and  
\* the accession number will be preserved.  
\* 1 231461: contig of 231461 bp in length.  
Location/Qualifiers  
1..231461  
/organism="Rattus norvegicus"  
/mol\_type="genomic DNA"  
/db\_xref="taxon:10116"  
/clone="CH230-11L24"  
1..1012  
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clone\_end:T7"  
2038..2914  
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site:ECORI  
228529..229317  
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## FEATURES

## source

## misc\_feature

## misc\_feature

## misc\_feature

## misc\_feature

## ORIGIN

Query Match 20.1%; Score 45.2; DB 2; Length 231461;  
Best Local Similarity 53.2%; Pred. No. 1;  
Matches 118; Conservative 0; Mismatches 103; Indels 1; Gaps 1;  
Qy 3 ATGTGAGTTGTTTCAGGCATATAAATTTGAATAAATATGAGCTCCATCATATGCTAT 62  
Db 150885 ATGAGATTTTAAATAGCATCAAGTATTAACCAATATTTTTCAGTTTATAGAA 150944  
Qy 63 ATTGGTTTTACCTTCAGAAGAATATTTAGTTTCACTCAGGTTTTTCAAAGCTACGCTGC 122  
Db 150945 GTGATGTATAATTTTAAATGTAACAATATTTTGTTCACCATTTGTCTAAGGAAGATTCT 151004  
Qy 123 CCCAAAAAGCAACAAACAAAAACAAACCTTTTAAAGATTGATGGTACTCATTT 182



humquery@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk  
 On Jan 17, 2003 this sequence version replaced gi:27764086.  
 Sequence from the Mouse Genome Sequencing Consortium whole genome  
 shotgun may have been used to confirm this sequence. Sequence data  
 from the whole genome shotgun alone has only been used where it has  
 a phred quality of at least 30.

----- Genome Center  
 Center: Wellcome Trust Sanger Institute  
 Center code: SC  
 Web site: <http://www.sanger.ac.uk>  
 Contact: [humquery@sanger.ac.uk](mailto:humquery@sanger.ac.uk)  
 -----

During sequence assembly data is compared from overlapping clones.  
 Where differences are found these are annotated as variations  
 together with a note of the overlapping clone name. Note that the  
 variation annotation may not be found in the sequence submission  
 corresponding to the overlapping clone, as we submit sequences with  
 only a small overlap as described above.

This sequence was finished as follows unless otherwise noted: all  
 regions were either double-stranded or sequenced with an alternate  
 chemistry or covered by high quality data (i.e., phred quality >=  
 30); an attempt was made to resolve all sequencing problems, such  
 as compressions and repeats; all regions were covered by at least  
 one plasmid subclone or more than one M13 subclone; and the  
 assembly was confirmed by restriction digest, except on the rare  
 occasion of the clone being a YAC.

The following abbreviations are used to associate primary accession  
 numbers given in the feature table with their source databases:  
 Em, EMBL; Sw, SWISSPROT; Tr, TrEMBL; Wp, WORMPEP; Information  
 on the WORMPEP database can be found at  
[http://www.sanger.ac.uk/Projects/C\\_elegans/wormpep](http://www.sanger.ac.uk/Projects/C_elegans/wormpep) RP23-324H1 is  
 from the RPCI-23 Mouse BAC Library

constructed by the group of Pieter de Jong.

For further details see <http://www.chori.org/bacpac/home.htm>

VECTOR: pBAC3.6.

Location/Qualifiers

1. 147727  
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 /mol\_type="genomic DNA"  
 /db\_xref="taxon:10090"  
 /chromosome="2"  
 /clone="RP23-324H1"  
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## ORIGIN

Query Match 19.7%; Score 44.4; DB 10; Length 147727;  
 Best Local Similarity 62.7%; Pred. No. 1.7;  
 Matches 69; Conservative 0; Mismatches 41; Indels 0; Gaps 0;

QY 74 CTTCAGAGAATATTAGTTTCACCTCAGGTTTTTCAAGCTACGCTGCCCCCAAAAC 133  
 Db 87490 CTACAAAGTGAGTTCAGGACCCTAGGCTATACAGAGAACACTGCTCGAAAAAAC 87431  
 QY 134 GAACAAACAAAAACACCTTTTAAAGATTGATGCTACTCTTTG 183  
 Db 87430 AAAACAAAAACAAACAAACATTCATTAGGTGGATTCTAAGAATTG 87381

## RESULT 14

AC104893/c  
 LOCUS AC104893 148750 bp DNA linear HTG 16-JUL-2003  
 DEFINITION HTG; HTGS\_PHASE2; HTGS\_DRAFT; HTGS\_FULLTOP.  
 pieces.  
 Mus musculus

AC104893  
 AC104893.4 GI:328113580

VERSION HTG; HTGS\_PHASE2; HTGS\_DRAFT; HTGS\_FULLTOP.

KEYWORDS Mus musculus (house mouse)

SOURCE Mus musculus

ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

1 (bases 1 to 148750)

REFERENCE Birren,B., Nusbaum,C. and Lander,E.

AUTHORS

TITLE Mus musculus, clone RP23-288O15

## JOURNAL

## REFERENCE

## AUTHORS

## Unpublished

## 2 (bases 1 to 148750)

Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N.,  
 Anderson,S., Barna,N., Bastien,V., Boguslavskiy,L., Boukhalter,B.,  
 Brown,A., Camarata,J., Campiano,A., Chang,J., Chazaro,B.,  
 Collier,Y., Colangelo,M., Collins,S., Collymore,A., Cook,A.,  
 Cooke,P., DeArelano,K., Dewar,K., Diaz,J.S., Dodge,S., Faro,S.,  
 Ferreira,P., FitzHugh,W., Gage,D., Galagan,J., Gardyna,S.,  
 Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N.,  
 Hagos,B., Heaford,A., Horton,L., Hulme,W., Iliev,I., Johnson,R.,  
 Jones,C., Kamat,A., Karatas,A., Kells,C., LaRoque,K.,  
 Lamazares,R., Landers,T., Lehoczy,J., Levine,R., Liu,G.,  
 Maclean,C., MacDonald,P., Major,J., Margis,N., Matthews,C.,  
 McCarthy,M., McEwan,P., McKernan,K., McPheeters,R., Meldrum,J.,  
 Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C.,  
 Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neil,D.,  
 Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V.,  
 Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P.,  
 Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S., Schupback,R.,  
 Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N.,  
 Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,  
 Topham,K., Travers,M., Travis,N., Trigilio,J., Vassiliev,H.,  
 Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,  
 Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

## Direct Submission

Submitted (22-DBC-2001) Whitehead Institute/MIT Center for Genome  
 Research, 320 Charles Street, Cambridge, MA 02141, USA

## 3 (bases 1 to 148750)

Birren,B., Nusbaum,C., Lander,E., Abouelleil,A., Allen,N.,  
 Anderson,M., Arachchi,H.M., Barna,N., Bastien,V., Bloom,T.,  
 Boguslavskiy,L., Boukhalter,B., Camarata,J., Chang,J., Choepe,Y.,  
 Collymore,A., Cook,A., Cooke,P., Corum,B., DeArelano,K.,  
 Diaz,J.S., Dodge,S., Dooley,K., Dorris,L., Erickson,J., Faro,S.,  
 Ferreira,P., FitzGerald,M., Gage,D., Galagan,J., Gardyna,S.,  
 Graham,L., Grand-Pierre,N., Hafez,N., Hagopian,D., Hagos,B.,  
 Hall,J., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C.,  
 Kamat,A., Karatas,A., Kells,C., Landers,T., Levine,R.,  
 Lindblad-Toh,K., Liu,X., Lui,A., Mabbitt,R., Maclean,C.,  
 MacDonald,P., Major,J., Manning,J., Matthews,C., McCarthy,M.,  
 Meldrum,J., Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J.,  
 Nguyen,C., Nicol,R., Norbu,C., O'Connor,T., O'Donnell,P.,  
 O'Neil,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N.,  
 Rachupka,A., Ramasamy,U., Raymond,C., Retta,R., Rise,C., Rogov,P.,  
 Roman,J., Schauer,S., Schupback,R., Seaman,S., Severy,P., Smith,C.,  
 Spencer,B., Stange-Thomann,N., Stojanovic,N., Stubbs,M.,  
 Talamas,J., Tesfaye,S., Theodore,J., Topham,K., Travers,M.,  
 Vassiliev,H., Venkataraman,V.S., Viel,R., Vo,A., Wilson,B., Wu,X.,  
 Wyman,D., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

## Direct Submission

Submitted (16-JUL-2003) Whitehead Institute/MIT Center for Genome  
 Research, 320 Charles Street, Cambridge, MA 02141, USA

On Jul 16, 2003 this sequence version replaced gi:20043160.

All repeats were identified using RepeatMasker:

Smit, A.F.A. & Green, P. (1996-1997)

-----ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBR

Web site: <http://www-seq.wi.mit.edu>

Contact: [sequence\\_submissions@genome.wi.mit.edu](mailto:sequence_submissions@genome.wi.mit.edu)

----- Project Information

Center project name: L19047

Center clone name: 288 O.15

----- Summary Statistics

Sequencing vector: Plasmid; n/a; 100% of reads

Chemistry: Dye-terminator Big Dye; 100% of reads

Assembly program: Phrap; version 0.960731

Consensus quality: 147390 bases at least Q40

Consensus quality: 147957 bases at least Q40

Consensus quality: 148226 bases at least Q20

Insert size: 152000; agarose-fp

Insert size: 148350; sum-of-contigs

Quality coverage: 9.7 in Q20 bases; agarose-fp







GenCore version 5.1.6  
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 27, 2004, 17:01:40 ; Search time 407 Seconds  
(without alignments)  
2348.512 Million cell updates/sec

Title: US-09-765-231A-58  
Perfect score: 225  
Sequence: 1 tgatgggaagtgttcagg.....attaggaattttttttttt 225

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 3373863 seqs, 212409041 residues

Total number of hits satisfying chosen parameters: 6747726

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database :

N Geneseq 29Jan04:\*  
1: Geneseqn1980s:\*  
2: Geneseqn1990s:\*  
3: Geneseqn2000s:\*  
4: Geneseqn2001as:\*  
5: Geneseqn2001bs:\*  
6: Geneseqn2002s:\*  
7: Geneseqn2003as:\*  
8: Geneseqn2003bs:\*  
9: Geneseqn2003cs:\*  
10: Geneseqn2004s:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	225	100.0	225	4	Aah23128 Osteoarth
2	225	100.0	320	7	Aca04823 CDNA enco
3	48.4	21.5	110000	2	Continuation (2 of
c 4	41	18.2	5518	6	Abk28306 DNA trans
c 5	40	17.8	6809	6	Abk31314 Signal tr
c 6	40	17.8	6809	6	Abk31314 Signal tr
c 7	40	17.8	6809	6	Abk31314 Signal tr
c 8	40	17.8	6809	6	Abk31314 Signal tr
c 9	38.2	17.0	110000	6	Abn80174 Human gen
c 10	38	16.9	12393	6	Continuation (2 of
c 11	37.8	16.8	96588	8	Abk33263 Human che
c 12	37.8	16.8	96588	8	Abk33263 Human che
c 13	37.8	16.8	96588	9	Abk33263 Human che
c 14	37.4	16.6	2270	2	Abk33263 Human che
c 15	37.2	16.5	778	4	Abk33263 Human che
c 16	36.6	16.3	2705	2	Abk33263 Human che
c 17	36.4	16.2	345	6	Abk33263 Human che
c 18	36.4	16.2	1479	6	Abk33263 Human che
c 19	36.4	16.2	7924	6	Abk33263 Human che
c 20	36.4	16.2	7924	6	Abk33263 Human che
c 21	36.4	16.2	40388	4	Abk33263 Human che
c 22	36.2	16.1	5474	6	Abk33263 Human che
c 23	36	16.0	5153	2	Aat30347 Human YAP

24	36	16.0	5153	9	ADD14716
c 25	36	16.0	6725	6	ABL33208 Human imm
c 26	36	16.0	6725	6	ABL34554 Human met
c 27	35.8	15.9	441	8	AD81862 Human CDN
c 28	35.8	15.9	502	6	ABQ88918 Human pro
c 29	35.8	15.9	3002	4	ABL15414 Drosophil
c 30	35.8	15.9	7847	6	ABL34188 Human imm
c 31	35.6	15.8	6163	6	ABN80119 Human che
c 32	35.4	15.7	932	4	AAK88641 Human dig
c 33	35.4	15.7	113515	6	ABL34175 Human imm
c 34	35.2	15.6	234	7	ABX54525 Bovine ES
c 35	35.2	15.6	557	4	AAI18969 Human bre
c 36	35.2	15.6	8038	9	ADB85272 Mouse vit
c 37	35.2	15.6	11049	6	ABL32668 Human imm
c 38	35.2	15.6	11049	6	ABL92218 Chemical
c 39	35.2	15.6	11049	6	ABL92218 Chemical
c 40	35.2	15.6	34319	8	ADA13460 Mouse rho
c 41	35	15.6	332	4	AAK55869 Human imm
c 42	35	15.6	1132	5	ABV28660 Human pro
c 43	35	15.6	1132	5	ABV22830 Human pro
c 44	35	15.6	2000	7	ADA71938 Rice gene
c 45	35	15.6	3541	2	AAV07076 CDNA enco

## ALIGNMENTS

### RESULT 1

AAH23128  
ID AAH23128 standard; DNA; 225 BP.

XX AC AAH23128;

DT 17-SEP-2001 (first entry)

XX DE Osteoarthritis tissue-derived nucleic acid sequence #58.

XX KW Osteoarthritis; infectious disorder; Crohn's disease; sepsis; human;  
wound healing; osteopathic; anti-arthritis; anti-inflammatory; vulnery;  
antibacterial; antiallergic; ds.

XX OS Homo sapiens.

XX PN WO200153531-A2.

XX PD 26-JUL-2001.

PR 18-JAN-2001; 2001WO-US0000016.

XX 18-JAN-2000; 2000US-0176523P.

XX (PHAA ) PHARMACIA CORP.

XX Phippard D, Vasanthakamur G, Dotson S, Ma X;

XX WPI; 2001-451914/48.

PT Substantially purified protein, polypeptide or their fragments, used to  
identify a biologically active compound or composition and treat  
mammalian osteoarthritis.

XX Claim 1; Page 137; 144pp; English.

XX Sequences AAH23071-23152 represent nucleic acid sequences derived from  
osteoarthritis tissues. The sequences are useful as probes and for the  
diagnosis or prognosis of mammalian osteoarthritis. The polynucleotides  
and polypeptides of the invention are useful for generating diagnostic  
reagents, as targets for small molecule drug development, generation of  
therapeutics, and cloning genes. Specific antibodies are used to generate  
enzyme linked immunosorbent assays for detection of osteoarthritis. The  
invented molecules can be used to treat osteoarthritis or to analyse the  
disease-modifying activity of osteoarthritis drugs. Other disorders  
treatable using the nucleic acid sequences include atopic, inflammatory



QY 67 GTTTTACCTTCAGAGATATTTAGTTCCTCAGGTTTTCAGAGTACGCTGTCCCCC 126  
 Db 51463 TCCTTATCAATAGCTCTCTAAATACTTCGTAATTTTAAACACTCCGGAGTTGTCT 51522  
 QY 127 AAAAAACGAAACAAAAACAAAAACAACTTTTAAAGAGTTGATGGCTACTCATTTGATC 186  
 Db 51523 AAAAAATCTATAAAATCAATATTACTCTTTTCCCACTCTTTAATTTGTTTTTATC 51582  
 QY 187 TGCTCTCTCTCTGAATCAATAGGAATTT 216  
 Db 51583 TTACCCAAATCCCACTATTAGGAATTT 51612

RESULT 4  
 ABK28306/c  
 ID ABK28306 standard; DNA; 5518 BP.

XX AC ABK28306;

XX DT 23-APR-2002. (first entry)

XX DE DNA transcription associated complementary genomic DNA #90.

XX KW DNA transcription associated gene; peptide nucleic acid; PNA-oligomer;  
 KW PNA; cytosine methylation state; SNP; retroviral infection; gene; ds;  
 KW single nucleotide polymorphism; adenosine deaminase deficiency; cancer;  
 KW viral infection; Sezary syndrome; haematological disorder; tuberculosis;  
 KW immunological disorder; Werner syndrome; developmental disorder;  
 KW psoriasis; Rieger's syndrome; neurological disorder; erythropoiesis;  
 KW neurodegenerative disorder; Waardenburg syndrome; Niemann-Pick disease;  
 KW myelodysplastic syndrome; myocardial infarction; hypertension; arthritis;  
 KW angiodysplasia; congenital heart disease; HDR syndrome; gene therapy;  
 KW polyglutamine disorder; solid tumour.

XX OS Unidentified.

XX PN WO200192565-A2.

XX PD 06-DEC-2001.

XX PF 06-APR-2001; 2001WO-EP003973.

XX PR 06-APR-2000; 2000DE-01019058.

XX PR 07-APR-2000; 2000DE-01019173.

XX PR 30-JUN-2000; 2000DE-01032529.

XX PR 01-SEP-2000; 2000DE-01043826.

XX PA (EPIG-) EPIGENOMICS AG.

XX PI Olek A, Piepenbrock C, Berlin K;

XX DR WPI; 2002-090046/12.

XX PT New nucleic acids or oligomers, useful for diagnosing or treating  
 PT diseases associated with DNA transcription, e.g. immunological disorders,  
 PT Werner syndrome, psoriasis, myocardial infarction, solid tumors or  
 PT cancer.

XX PS Claim 1; SEQ ID NO 180; 32pp; English.

XX CC The invention relates to a nucleic acid, which comprises a segment of the  
 CC chemically pretreated DNA of genes associated with DNA transcription from  
 CC one of 346 sequences, and an oligomer, in particular an oligonucleotide  
 CC or peptide nucleic acid (PNA)-oligomer that hybridises to or is identical  
 CC to the chemically pretreated DNA of genes associated with DNA  
 CC transcription. The set of oligomer probes are useful for detecting the  
 CC cytosine methylation state and/or single nucleotide polymorphisms (SNPs)  
 CC in a chemically pretreated genomic DNA. The nucleic acids are useful for  
 CC diagnosing or treating diseases associated with DNA transcription  
 CC (particularly with the methylation status), e.g. adenosine deaminase  
 CC deficiency, viral infection, retroviral infection, Sezary syndrome,  
 CC haematological disorders, immunological disorders, Werner syndrome,  
 CC tuberculosis, developmental disorders, psoriasis, Rieger's syndrome,

CC neurological disorders, neurodegenerative disorders, Waardenburg  
 CC syndrome, Niemann-Pick disease, myelodysplastic syndrome, myocardi  
 CC infarction, hypertension, angiodysplasia, erythropoiesis, congenital heart  
 CC disease, HDR syndrome, arthritis, polyglutamine disorders, solid tumours  
 CC or cancer. Sequences ABK28127-ABK28472 represent DNA transcription  
 CC associated genomic DNA molecules of the invention. Note: The sequence  
 CC data for this patent did not form part of the printed specification but  
 CC was obtained in electronic format directly from the European Patent  
 CC Office

XX SQ Sequence 5518 BP; 1899 A; 52 C; 1031 G; 2536 T; 0 U; 0 Other;

Query Match 18.2%; Score 41; DB 6; Length 5518;

Best Local Similarity 51.4%; Pred. No. 1.5;

Matches 95; Conservative 0; Mismatches 90; Indels 0; Gaps 0;

QY 7 TAAGTTGTTTCAGGCATATAAATTTGAAATAAATATGAGGCTCATGATATGCTATATTG 66

Db 4088 TATATTTTAAACAAAAATTTTCATTTAACTATAAACTCTCTATAAATATTACGAA 4029

QY 67 GTTTTACCTTCAGAGATATTTAGTTCCTCAGGTTTTCAGAGTACGCTGTCCCCC 126

Db 4028 TTACTTAAATTAATAATAATAATAATAATAATAATAATAATAATAATAATAATAATA 3969

QY 127 AAAAAACGAAACAAAAACAAAAACAACTTTTAAAGAGTTGATGGCTACTCATTTGATC 186

Db 3968 AATAAAAAATCTAATAATAATAATAATAATAATAATAATAATAATAATAATAATAATA 3909

QY 187 TGCTCT 191

Db 3908 TAACT 3904

RESULT 5

ID ABK31314/c

XX ABK31314 standard; DNA; 6809 BP.

XX AC ABK31314;

XX DT 23-APR-2002 (first entry)

XX DE Signal transduction associated gene modified DNA #79.

XX KW Human; signal transduction associated gene; cytosine methylation state;  
 KW CpG island; signal transduction associated disease; solid tumour; cancer;  
 KW antitumour; cytostatic; mutant; ds.

XX OS Homo sapiens.

XX OS Synthetic.

XX PN WO200200926-A2.

XX PD 03-JAN-2002.

XX PF 29-JUN-2001; 2001WO-EP007472.

XX PR 30-JUN-2000; 2000DE-01032529.

XX PR 01-SEP-2000; 2000DE-01043826.

XX PA (EPIG-) EPIGENOMICS AG.

XX PI Olek A, Piepenbrock C, Berlin K;

XX DR WPI; 2002-147896/19.

XX PT Oligonucleotide for diagnosis and therapy of diseases associated with  
 PT signal transduction e.g. cancer, comprises chemically modified genomic  
 PT sequences of genes associated with signal transduction.  
 XX Claim 1; SEQ ID NO 157; 24pp; English.

XX CC The present invention relates to chemically modified DNA sequences of  
 CC signal transduction associated genes. The DNA sequences are chemically

CC modified using a solution of bisulphite, hydrogen sulphite or disulphite.  
CC Also disclosed are oligonucleotides and/or DNA oligomers for detecting  
CC the cytosine methylation state (CpG islands) of these genes, and a method  
CC for the diagnosis and/or therapy of genetic and epigenetic parameters of  
CC genes associated with signal transduction. The genomic DNA can be  
CC obtained from cells or cellular components which contain DNA, e.g. cell  
CC lines, biopsies, blood, sputum, stool, urine, cerebral-spinal fluid,  
CC tissue embedded in paraffin such as tissue from eyes, intestine, kidney,  
CC brain, heart, prostate, lung, breast or liver, histologic object slides,  
CC and all their possible combinations. The sequences of the invention are  
CC useful for the diagnosis and therapy of diseases associated with signal  
CC transduction e.g. solid tumours and cancer. ABK31158-ABK31545 represent  
CC chemically pretreated genomic DNA sequences of different genes associated  
CC with signal transduction, or their complementary sequences. Note: The  
CC sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from the  
CC European Patent Office  
XX  
XX  
SQ Sequence 6809 BP; 1792 A; 270 C; 1625 G; 3122 T; 0 U; 0 Other;

Query Match 17.8%; Score 40; DB 6; Length 6809;  
Best Local Similarity 54.9%; Pred. No. 2.7;  
Matches 79; Conservative 0; Mismatches 65; Indels 0; Gaps 0;  
Qy 17 CAGGCATAAATTGGAATAAAATATGAGGCTCCATGATATGCTATATTGCTTTTACCTT 76  
Db 1384 CACAAATAAAACCAATATATATCGTACACACTCTATCATATTTCTTTAATAT 1325  
Qy 77 CAGAAGATATTTAGTTTCACTCAGGTTTTTCAAAGCTACGCTGTCCCCCAAAAACGAA 136  
Db 1324 ACAAAACTATTTAAATTAATCCGCTCATCAATTTTTCGTTTACTACAAATTACTTT 1265  
Qy 137 ACAAAACAAAACCAACCTTTT 160  
Db 1264 TCAAACTTAATCATATAATCTTT 1241

RESULT 6  
ABL70557/c  
ID ABL70557 standard; DNA; 6809 BP.  
XX  
AC ABL70557;  
XX  
DT 01-JUL-2002 (first entry)  
XX  
DE Chemically treated cell signalling DNA sequence#224.  
XX  
KW Cell signalling; cytosine methylation; cell signalling disease; cancer;  
KW tumour; cytostatic; ds.  
XX  
OS Unidentified.  
XX  
FN WO200202807-A2.  
XX  
PD 10-JAN-2002.  
XX  
PF 29-JUN-2001; 2001WO-EP007471.  
XX  
XX 30-JUN-2000; 2000DE-01032529.  
PR 01-SEP-2000; 2000DE-01043826.  
XX  
PA (EPIG-) EPIGENOMICS AG.  
XX  
FI Olek A, Piepenbrock C, Berlin K;  
XX  
DR WPI; 2002-154758/20.  
XX  
PT Nucleic acid, useful for diagnosis and therapy of diseases associated  
PT with cell signaling e.g. cancer, comprises chemically modified genomic  
PT sequences of genes associated with cell signaling.  
XX  
PS Claim 1; SEQ ID NO 447; 24pp + Sequence Listing; English.  
XX

CC The invention relates to a nucleic acid comprising a sequence of at least  
CC 18 bases of a segment of chemically pretreated DNA of genes associated  
CC with cell signalling. The activity of the modified sequences of the  
CC invention may be described as cytostatic. The object of the invention is  
CC to provide the chemically modified DNA of genes associated with cell  
CC signalling, as well as oligonucleotides and/or DNA oligomers for  
CC detecting cytosine methylations, as well as a method which is  
CC particularly suitable for the diagnosis and/or therapy of genetic and  
CC epigenetic parameters of genes associated with cell signalling. The  
CC chemically modified DNA provided by the invention is useful for diagnosis  
CC given in records ABL70111-ABL70626 represent chemically pre-treated  
CC genomic DNA's of genes associated with cell signalling. Note: The  
CC sequence data for this patent is not represented in the printed  
CC specification, but is based on sequence information supplied by the  
CC European Patent Office  
XX  
XX  
SQ Sequence 6809 BP; 1792 A; 270 C; 1625 G; 3122 T; 0 U; 0 Other;

Query Match 17.8%; Score 40; DB 6; Length 6809;  
Best Local Similarity 54.9%; Pred. No. 2.7;  
Matches 79; Conservative 0; Mismatches 65; Indels 0; Gaps 0;  
Qy 17 CAGGCATAAATTGGAATAAAATATGAGGCTCCATGATATGCTATATTGCTTTTACCTT 76  
Db 1384 CACAAATAAAACCAATATATATCGTACACACTCTATCATATAATTTCTTTAATAT 1325  
Qy 77 CAGAAGATATTTAGTTTCACTCAGGTTTTTCAAAGCTACGCTGTCCCCCAAAAACGAA 136  
Db 1324 ACAAAACTATTTAAATTAATCCGCTCATCAATTTTTCGTTTACTACAAATTACTTT 1265  
Qy 137 ACAAAACAAAACCAACCTTTT 160  
Db 1264 TCAAACTTAATCATATAATCTTT 1241

RESULT 7  
AAS61214/c  
ID AAS61214 standard; DNA; 6809 BP.  
XX  
AC AAS61214;  
XX  
DT 29-JAN-2002 (first entry)  
XX  
DE Human gene regulation-associated gene oligonucleotide #169.  
XX  
KW Human; Gene regulation-associated gene; severe combined immunodeficiency;  
KW cardiac damage; inflammatory response; Haemophilia; Werner syndrome;  
KW asthma; HDR syndrome; congenital heart defect; Saethre-Chotzen syndrome;  
KW renal disease; Preeclampsia; cardiac allograft vascular disease;  
KW colorectal cancer; thyroid cancer; oesophageal cancer; ds; tumour;  
KW immunostimulant; cardiant; antiinflammatory; coagulant; antiasthmatic;  
KW nephrotropic; gynecological; anti-tumour; immunosuppressive; cytostatic.  
XX  
OS Homo sapiens.  
XX  
FN WO200177375-A2.  
XX  
PD 18-OCT-2001.  
XX  
PF 06-APR-2001; 2001WO-EP003968.  
XX  
XX 06-APR-2000; 2000DE-01019058.  
PR 07-APR-2000; 2000DE-01019173.  
PR 30-JUN-2000; 2000DE-01032529.  
PR 01-SEP-2000; 2000DE-01043826.  
XX  
PA (EPIG-) EPIGENOMICS AG.  
XX  
FI Olek A, Piepenbrock C, Berlin K;  
XX  
DR WPI; 2002-017470/02.  
XX



Db 98242 TAAATGTTGTTTATTAATAATACCTTTTCATCTAAATAAACAATACAAAATAAAAAA 98183

Qy 152 AACCTTTTAA 162

Db 98182 AAGATTTTTTA 98172

RESULT 10

ABL33263/c

ID ABL33263 standard; DNA; 12393 BP.

XX AC ABL33263;

XX DT 26-MAR-2002 (first entry)

XX DE Human immune system associated gene SEQ ID NO: 1236.

XX KW Human; immune system disease; cytosine methylation; antiasthmatic;

XX KW antiarteriosclerotic; antianemic; cytostatic; nootropic;

XX KW neuroprotective; anti-HIV; anticonvulsant; ophthalmological;

XX KW antirheumatic; antiarthritic; antididiabetic; antiporiatic;

XX KW antiinflammatory; cancer; eye disease; arteriosclerosis; anaemia;

XX KW acute myeloid leukaemia; Alzheimer's disease; AIDS; epilepsy;

XX KW neurofibromatosis; rheumatoid arthritis; psoriasis; bowel disease; gene;

XX KW ds.

XX OS Homo sapiens.

XX PN WO200200928-A2.

XX PD 03-JAN-2002.

XX PF 02-JUL-2001; 2001WO-EP007537.

XX PR 30-JUN-2000; 2000DE-01032529.

XX PR 01-SEP-2000; 2000DE-01043826.

XX PA (EPIG-) EPIGENOMICS AG.

XX PI Olek A, Piepenbrock C, Berlin K;

XX DR WPI; 2002-130909/17.

XX PT Nucleic acid comprising fragment of chemically modified gene, useful for

XX PT diagnosis and treatment of diseases associated with abnormal cytosine

XX PT methylation.

XX PS Claim 1; SEQ ID NO 1236; 32pp + Sequence Listing; German.

XX CC The present invention provides a number of human immune system associated

XX CC genes which are modified by the methylation of cytosines. The sequences

XX CC can be used in the diagnosis and treatment of immune system disorders,

XX CC including eye diseases such as retinopathy, neovascular glaucoma and

XX CC macular degeneration, arteriosclerosis, anaemia, cancer, acute myeloid

XX CC leukaemia, Alzheimer's disease, AIDS, epilepsy, neurofibromatosis,

XX CC rheumatoid arthritis, psoriasis and inflammatory/ulcerative bowel

XX CC diseases. The present sequence is a gene of the invention

XX CC

SQ Sequence 12393 BP; 3484 A; 219 C; 2406 G; 6282 T; 0 U; 2 Other;

Query Match 16.9%; Score 38; DB 6; Length 12393;

Best Local Similarity 49.5%; Pred. No. 9;

Matches 98; Conservative 0; Mismatches 100; Indels 0; Gaps 0;

Qy 22 ATAAATTTGAATAAATATGAGGCTCCATGATATGCTATATGTTTACTTCAGAA 81

Db 6387 AAAAAAATAAAAAAATTTAAAAATAATTTAAAAATAAATTTTCTCTTAAA 6328

Qy 82 GAATATTAGTTTCACTCAGGTTTTCAGAGTCGCTGCCCAAAAAACGAACAA 141

Db 6327 AAAACATAATTTTATTACGATTTTAAAAAATCAAAATACTCGATAAATAATA 6269

Qy 142 ACAAAAAACAACCTTTTAAAGATTGATGGCTACTCAATTGATCTGCTCTCTCTGTA 201

Db 6267 TAAATAAATAAATAATATCTCAATTCTAACTTAATTCTCCGCTCTCTCAATAACCTTA 6208

Qy 202 ATCAATTAGGAATTTTT 219

Db 6207 AACAAATTTCTATCAATT 6190

RESULT 11

ADA03026/c

ID ADA03026 standard; DNA; 96588 BP.

XX AC ADA03026;

XX DT 06-NOV-2003 (first entry)

XX DE Human MBNL carcinoma associated gene, SEQ ID NO:1544.

XX KW Human; carcinoma associated; oncogene; carcinoma; cancer; breast;

XX KW prostate; lymphoma; leukaemia; cytostatic; gene therapy; drug screening;

XX KW gene; ds.

XX OS Homo sapiens.

XX PN WO2003057146-A2.

XX PD 17-JUL-2003.

XX PF 26-DEC-2002; 2002WO-US041414.

XX PR 26-DEC-2001; 2001US-00035832.

XX PA (SAGR-) SAGRES DISCOVERY.

XX PI Morris DW;

XX DR WPI; 2003-587068/55.

XX PT New recombinant nucleic acid encoding carcinoma associated protein,

XX PT useful for preparing compositions for treating carcinomas.

XX PS Claim 1; SEQ ID NO 1544; 245pp; English.

XX CC The invention relates to recombinant carcinoma associated (CA) nucleic

XX CC acid sequences from mouse and human (ADA01482-ADA03094), and to

XX CC recombinant carcinoma associated proteins (CAP) encoded by them. The

XX CC invention also encompasses expression vectors and host cells comprising a

XX CC CA nucleic acid, a polypeptide (especially an antibody) that specifically

XX CC binds to the protein, and a biochip comprising CA nucleic acid or

XX CC fragments thereof. The sequences of the invention were identified using

XX CC oncogenic retroviruses, which insert into the genome of the host organism

XX CC at random. Many of these do not carry transduced host oncogenes or

XX CC pathogenic trans-acting viral genes, meaning that cancer incidence is a

XX CC direct consequence of the effects of proviral integration into host

XX CC protooncogenes. The CA nucleic acid sequences can be used to diagnose

XX CC carcinoma (especially breast cancer, prostate cancer, lymphoma or

XX CC leukaemia) or a propensity to carcinoma by determination of the sequence

XX CC of a CA gene, or by determination of CA gene expression in particular

XX CC tissues. CA nucleic acids, proteins and antibodies are also useful as

XX CC therapeutic agents and in screening and evaluating drug candidates. The

XX CC present sequence represents a specifically claimed human CA nucleic acid

XX CC sequence of the invention. Note: The complete sequence data for this

XX CC patent did not form part of the printed specification, but was obtained

XX CC in electronic format directly from WIPO at

XX CC ftp.wipo.int/pub/published\_pct\_sequences.

XX CC

SQ Sequence 96588 BP; 29654 A; 16428 C; 18069 G; 32437 T; 0 U; 0 Other;

Query Match 16.8%; Score 37.8; DB 8; Length 96588;

Best Local Similarity 51.5%; Pred. No. 13;

Matches 87; Conservative 0; Mismatches 82; Indels 0; Gaps 0;

QY 8 AAGTTGTTTCAGGCATATAAATTTGAATAATATGAGGCTCCATGATGCTATATTGG 67  
 Db 61750 ATGTTGAGCATGTGAATTTTATATAATTTAAATTTGCTTCCATGCTCAATTTATTAGTA 61691  
 QY 68 TTTTACCTTCAGAGAAATATTAGTTTTCACCTCAGGTTTTCAGAGCTACGCTGCCCCCA 127  
 Db 61690 AATTTTATAAATAGCATATGATACACATGCAATTTTTCAGAGCTGCTGTTGCCAGCC 61631  
 QY 128 AAAACAGCAACAAACAAACAAACAAACCTTTTAAAGATTGATGCTGCTAC 176  
 Db 61630 AAAAAAAAAAAAAAAAAAAAAAAAAAAGTATAGAACTTTTAATGCCAC 61582

## RESULT 12

ADB72764/c

ID ADB72764 standard; DNA; 96588 BP.

XX AC  
 XX ADB72764;

04-DEC-2003 (first entry)

Human MBNL gene.

XX human; ds; cytostatic; gene therapy; vaccine; carcinoma; lymphomas;  
 KW cancer; neoplasm; adenocarcinoma; sarcoma; gene.

XX Homo sapiens.

XX WO2003008583-A2.

XX 30-JAN-2003.

XX 26-DEC-2001; 2001WO-US051291.

XX 02-MAR-2001; 2001US-00798586.

XX 23-OCT-2001; 2001US-00004113.

XX 08-NOV-2001; 2001US-00052482.

XX 30-NOV-2001; 2001US-00997722.

XX 20-DEC-2001; 2001US-00034650.

XX (SAGR-) SAGRES DISCOVERY.

XX Morris DW, Engelhard EK;

XX WPI; 2003-239337/23.

XX New recombinant nucleic acid, useful for treating carcinomas, lymphomas,  
 PT cancers, neoplasm, adenocarcinoma, or sarcomas.

XX Claim 1; SEQ ID NO 592; 2304pp; English.

XX The invention relates to a novel recombinant nucleic acid comprising a  
 CC nucleotide sequence selected from any of the 660 sequences fully defined  
 CC in the specification. A polynucleotide of the invention has cytostatic  
 CC activity, and may have a use in gene therapy, or in a vaccine. The  
 CC recombinant nucleic acids and polypeptides are useful for treating  
 CC carcinomas, e.g. lymphomas, cancers, neoplasm, adenocarcinoma, and  
 CC sarcomas. The present sequence represents a human gene of the invention.

XX Sequence 96588 BP; 29654 A; 16428 C; 18069 G; 32437 T; 0 U; 0 Other;

Query Match

Best Local Similarity 16.8%; Score 37.8; DB 9; Length 96588;

Matches 87; Conservative 0; Mismatches 82; Indels 0; Gaps 0;

QY 8 AAGTTGTTTCAGGCATATAAATTTGAATAATATGAGGCTCCATGATGCTATATTGG 67  
 Db 61750 ATGTTGAGCATGTGAATTTTATATAATTTAAATTTGCTTCCATGCTCAATTTATTAGTA 61691

QY 68 TTTTACCTTCAGAGAAATATTAGTTTTCACCTCAGGTTTTCAGAGCTACGCTGCCCCCA 127  
 Db 61690 AATTTTATAAATAGCATATGATACACATGCAATTTTTCAGAGCTGCTGTTGCCAGCC 61631

QY 128 AAAACAGCAACAAACAAACAAACAAACCTTTTAAAGATTGATGCTGCTAC 176  
 Db 61630 AAAAAAAAAAAAAAAAAAAAAAAAAAAGTATAGAACTTTTAATGCCAC 61582

## RESULT 13

ADC85506/c

ID ADC85506 standard; DNA; 96588 BP.

XX AC  
 XX ADC85506;

01-JAN-2004 (first entry)

Human Mbnl genomic sequence.

XX Cytostatic; gene therapy; vaccine; cancer; carcinoma-associated gene; CA;  
 KW secreted; transmembrane; intracellular; ds.

XX Homo sapiens.

XX WO2003045230-A2.

XX 05-JUN-2003.

XX 02-DEC-2002; 2002WO-US038582.

XX 30-NOV-2001; 2001US-00997722.

XX (SAGR-) SAGRES DISCOVERY.

XX Morris DW, Engelhard EK;

XX WPI; 2003-513603/48.

XX New recombinant nucleic acid comprising a nucleotide sequence of any of  
 PT the carcinoma-associated (CA) genes, useful for screening for drug  
 PT candidates for diagnosing or treating carcinomas.

XX Claim 1; SEQ ID NO 292; 983pp; English.

XX The invention relates to a recombinant nucleic acid comprising a  
 CC nucleotide sequence selected from any of the fully defined carcinoma-  
 CC associated (CA) genes from the 50 tables given in the specification. The  
 CC CA proteins are secreted, transmembrane or intracellular proteins. The  
 CC recombinant nucleic acids are useful for screening for drug candidates  
 CC for diagnosing or treating carcinomas. Sequences given in ADC85215-  
 CC ADC85514 represent CA genes of the invention.

XX Sequence 96588 BP; 29654 A; 16428 C; 18069 G; 32437 T; 0 U; 0 Other;

Query Match

Best Local Similarity 16.8%; Score 37.8; DB 9; Length 96588;

Matches 87; Conservative 0; Mismatches 82; Indels 0; Gaps 0;

QY 8 AAGTTGTTTCAGGCATATAAATTTGAATAATATGAGGCTCCATGATGCTATATTGG 67  
 Db 61750 ATGTTGAGCATGTGAATTTTATATAATTTAAATTTGCTTCCATGCTCAATTTATTAGTA 61691

QY 68 TTTTACCTTCAGAGAAATATTAGTTTTCACCTCAGGTTTTCAGAGCTACGCTGCCCCCA 127  
 Db 61690 AATTTTATAAATAGCATATGATACACATGCAATTTTTCAGAGCTGCTGTTGCCAGCC 61631

QY 128 AAAACAGCAACAAACAAACAAACAAACCTTTTAAAGATTGATGCTGCTAC 176  
 Db 61630 AAAAAAAAAAAAAAAAAAAAAAAAAAAGTATAGAACTTTTAATGCCAC 61582

## RESULT 14

AAK05715/c

ID AAK05715 standard; DNA; 2270 BP.

XX

XX AAK05715;

XX

DT 07-MAY-1999 (first entry)  
XX Human protein phosphatase (PROPHO) encoding DNA.  
DE  
XX  
XX Protein phosphatase; PROPHO; apoptosis; AIDS; Alzheimer's Disease;  
KW Acquired Immune Deficiency Syndrome; Parkinson's Disease; inflammation;  
KW cell proliferation; Addison's disease; allergy; anemia; cancer; bone;  
KW leukemia; breast; brain; human; ss.  
XX  
XX Homo sapiens.  
OS  
XX Key Location/Qualifiers  
FH 70..1509  
FT CDS /\*tag= a  
FT /product= "protein phosphatase (PROPHO) "  
XX  
XX W09856925-A1.  
XX  
XX 17-DEC-1998.  
PD  
XX 11-JUN-1998; 98WO-US011614.  
XX  
XX 11-JUN-1997; 97US-00873093.  
PR  
XX (INCY-) INCYTE PHARM INC.  
PA  
XX Bandman O, Goli SK, Lal P, Corley NC, Zhang H;  
PI  
XX WPI; 1999-080906/07.  
DR P-PSDB; AAW94283.  
XX  
XX New substantially purified human protein phosphatase (PROPHO) - useful in  
PT the diagnosis, prevention or treatment of inflammation, cancer, and  
PT disorders associated with apoptosis.  
XX  
XX Claim 5; Fig 1A-G; 73pp; English.  
XX  
XX This DNA encodes a human protein phosphatase (PROPHO). Host cells  
CC containing a vector comprising the PROPHO nucleic acid are used for the  
CC recombinant production of the protein. PROPHO forms a composition in the  
CC treatment or prevention of apoptosis-related disorders (e.g. Acquired  
CC Immune Deficiency Syndrome (AIDS), Alzheimer's Disease and Parkinson's  
CC Disease), and in the stimulation of cell proliferation. Antagonists of  
CC the protein are useful in treating inflammation (e.g. Addison's disease,  
CC allergies and anemia), and disorders associated with cell proliferation  
CC (including various cancers like leukemia, and cancers affecting bone,  
CC breast and brain). Complementary polynucleotides are useful in detecting  
CC polynucleotides that encode PROPHO, useful in the diagnosis of conditions  
CC associated with the expression of PROPHO, and in assays that detect  
CC activation or induction of various cancers. PROPHO is useful in producing  
CC antibodies or screening libraries of pharmaceutical agents in order to  
CC identify those that bind to PROPHO  
XX  
SQ Sequence 2270 BP; 693 A; 344 C; 515 G; 716 T; 0 U; 2 Other;  
Query Match 16.6%; Score 37.4; DB 2; Length 2270;  
Best Local Similarity 53.0%; Pred. No. 10;  
Matches 80; Conservative 0; Mismatches 71; Indels 0; Gaps 0;  
QY 29 TTGAATAAATATCAGGCTCCATGATGCTATATGTTTACCTTCAGAAATATT 88  
Db 2197 TTGACATACATTTTGTAGTGTAGTGTAGTGTATACAGAAATATACAAACCAAGCTGTAT 2138  
QY 89 TAGTTTCTACTAGGTTTTCRAAGCTACGCTCTCCCAAAACGAAACAAACAAA 148  
Db 2137 GAATAATACATAGGTTTTCRAAGTTATGTTTTCATAAGAAATACAGAAAGGAAAT 2078  
QY 149 AACAACTTTTAAAGAGTTGATGCTACTCA 179  
Db 2077 AGCCACACATCCAAATATCTCACAACTTCTAA 2047

RESULT 15

AAAL21862/c  
ID AAL21862 standard; cDNA; 778 BP.  
XX  
XX AAL21862;  
AC  
XX 07-DEC-2001 (first entry)  
DT  
XX Human breast cancer expressed polynucleotide 14319.  
DE  
XX Human; breast cancer; cell marker; cytostatic; ss.  
KW  
XX Homo sapiens.  
OS  
XX W0200151628-A2.  
PN  
XX 19-JUL-2001.  
PD  
XX 10-JAN-2001; 2001WO-US000798.  
PF  
XX 14-JAN-2000; 2000US-0176077P.  
PR  
XX 14-MAR-2000; 2000US-0189167P.  
PR  
XX 24-MAR-2000; 2000US-0192099P.  
PR  
XX 29-MAR-2000; 2000US-0193480P.  
PR  
XX 15-MAY-2000; 2000US-0205230P.  
PR  
XX 09-JUN-2000; 2000US-0211315P.  
PR  
XX 25-JUL-2000; 2000US-0220534P.  
XX (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.  
PA  
XX Lillie J, Xu Y, Wang Y, Steinmann K;  
PI  
XX WPI; 2001-451856/48.  
DR  
XX New peptide useful as a marker for the diagnosis of breast cancer.  
PT  
XX  
XX Claim 1; Page 2564-2565; 3695pp; English.  
XX  
XX The invention relates to human breast cancer expressed polynucleotides  
CC (AAL07544-AA26789) and methods of assessing whether a patient is  
CC afflicted with breast cancer by examining the correlation between the  
CC expression of certain markers and the cancerous state of breast cells.  
CC The polynucleotides and encoded polypeptides are potential markers for  
CC detecting, diagnosing, monitoring, characterizing treating and  
CC potentially preventing breast cancer. The polynucleotides and encoded  
CC polypeptides are also useful for isolating compounds with cytostatic  
CC activity  
XX  
SQ Sequence 778 BP; 220 A; 134 C; 180 G; 244 T; 0 U; 0 Other;  
Query Match 16.5%; Score 37.2; DB 4; Length 778;  
Best Local Similarity 55.4%; Pred. No. 9.8;  
Matches 72; Conservative 0; Mismatches 58; Indels 0; Gaps 0;  
QY 27 ATTGGAATAAATTATGAGGCTCCATGATGCTATATGTTTACCTTCAGAAATA 86  
Db 136 AATTTAGCTATATTTATCTTCTCAATCAAAATTTCTACTCAGAGGTAAGTAAAGAAAGT 77  
QY 87 TTTAGTTTCACTCAGGTTTTTCAAAGCTACGCTGTCCCAAAACGAAACCAACAAA 146  
Db 76 TATAGCAATCATATAAATGGACAAAAAGGTATCCCAAAAAAANAANAANA 17  
QY 147 AAAACAACCT 156  
Db 16 AAAAGTACCT 7  
Search completed: July 27, 2004, 18:19:56  
Job time : 411 secs